

GenCore version 5.1.6
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OM protein - nucleic search, using frame_plus_p2n model

Run on: May 4, 2005, 09:07:56 ; Search time 363.973 Seconds
(without alignments)
836.639 Million cell updates/sec

Title: US-09-017-715A-2_COPY_120_127
Perfect score: 41
Sequence: 1 EEAQSGSD 8

Scoring table:
BLOSUM62
Xgapop 10.0, Xgapext 0.5
Ygapop 10.0, Ygapext 0.5
Fgapop 6.0, Fgapext 7.0
Delop 6.0, Delext 7.0

Searched: 34239544 seqs, 19032134700 residues
Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Command line parameters:
-MODEL=frame+ p2n.model -DEV=xlh
-O=/cgr2_1/USPTO.spool.h/US090171715/runat_04052005_100744_25619/app_query.fasta_1.661
-DB=EST -OPMT=fastap -SUFFIX=rest -MINMATCH=0.1 -LOOPEXT=0 -LOOPEXT=0
-UNITS=bits -START=1 -END=1 -MATRIX=bl0sum62 -TRANS=human40.cdi -LIST=45
-LOCAL=GN=200 -THR_SCORE=pct -THR_MAX=100 -THR_MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTFMT=pro -NOR=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=200000000
-USRR=US090171715_@CGN_1_1_5334_@runat_04052005_100744_25619 -NCPU=6 -ICPU=3
-NO_MMAP -LARGEQUERY -NEG_SCORES=0 -WAIT -DSPBLOCK=100 -LONGCLOG
-DEV_TIMEOUT=120 -MAIN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -Fgapop=6
-Fgapext=7 -Ygapop=10 -Ygapext=0.5 -Delop=6 -Delext=7

Database :
EST:
1: gb_esc1:*
2: gb_esc2:*
3: gb_hic:*
4: gb_esc3:*
5: gb_esc4:*
6: gb_esc5:*
7: gb_esc6:*
8: gb_gss1:*
9: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	41	100.0	233	2	BE171313 RCI-HT054
2	41	100.0	301	1	AA055968 zf22a02.8
3	41	100.0	327	1	AA056035 zf22a02.r
4	41	100.0	333	4	BM748818 K-EST0023
5	41	100.0	352	1	A1468480 t957912.x
6	41	100.0	368	5	BU729743 UI-E-CK1
7	41	100.0	378	1	AA293803 zt56h04.8
8	41	100.0	387	1	AA722407 z983h10.8
9	41	100.0	393	1	AA394097 zt56h04.r

C	10	41	100.0	404	7	CF529590	UI-1-BC1P
C	11	41	100.0	407	1	AV683707	AV683707
C	12	41	100.0	408	1	AA946606	AA946606
C	13	41	100.0	412	1	A1139933	qae6b04.x
C	14	41	100.0	421	1	AV703171	AV703171
C	15	41	100.0	428	5	BX090816	BX090816
C	16	41	100.0	438	1	A1684600	wa84d12.x
C	17	41	100.0	438	1	A1936527	wd29a07.x
C	18	41	100.0	442	1	AA804675	0f44b01.8
C	19	41	100.0	451	4	BG826435	BG826435
C	20	41	100.0	472	4	BM704200	BM704200
C	21	41	100.0	473	6	CA421283	CA421283
C	22	41	100.0	474	4	BM706956	BM706956
C	23	41	100.0	480	4	BM668990	BM668990
C	24	41	100.0	504	5	BU728272	UI-E-CK0
C	25	41	100.0	509	2	BE299889	BE299889
C	26	41	100.0	510	2	BE298825	60119383
C	27	41	100.0	519	1	A1016464	0c78h06.8
C	28	41	100.0	538	4	BM695726	UI-E-CK1
C	29	41	100.0	544	4	BM655098	UI-E-CK1
C	30	41	100.0	553	1	AA872836	0h76f10.8
C	31	41	100.0	555	5	BX474500	DKFZP686D
C	32	41	100.0	555	7	CV028548	7090_Full1
C	33	41	100.0	558	4	B1548891	603189023
C	34	41	100.0	568	4	BG708703	602674249
C	35	41	100.0	578	5	BP212912	BP212912
C	36	41	100.0	582	5	BP197662	BP197662
C	37	41	100.0	583	5	BP200612	BP200612
C	38	41	100.0	583	5	BP346497	BP346497
C	39	41	100.0	584	1	AV708933	AV708933
C	40	41	100.0	584	5	BP201686	BP201686
C	41	41	100.0	588	4	BI755243	603023269
C	42	41	100.0	592	6	CA443299	UI-H-DH1
C	43	41	100.0	593	6	CA443166	UI-H-DH1
C	44	41	100.0	614	5	BU730570	UI-E-C11
C	45	41	100.0	617	1	AA633976	ac33e03.8

ALIGNMENTS

RESULT 1	BE171313	233 bp	mrna	linear	EST 21-JUN-2000
LOCUS	BE171313				
DEFINITION	RCI-HT0545-100300-012-g05 HT0545 Homo sapiens cDNA, mRNA sequence.				
ACCESSION	BE171313				
VERSION	BE171313.1	GI:8634039			
KEYWORDS	EST.				
SOURCE	Homo sapiens (human)				
ORGANISM	Homo sapiens				
REFERENCE	Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.				
AUTHORS	1 (bases 1 to 233) Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Britones,M.R., Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F., Goldman,G.H., Carvalho,A.F., Matsukuma,A., Bata,G.S., Simpson,D.H., Brumstein,A., deOliveira,P.S., Bucher,P., Jorgensen,C.V., O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and Simpson,A.J.				
TITLE	Shotgun sequencing of the human transcriptome with ORF expressed sequence tags				
JOURNAL	Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)				
MEDLINE	20202663				
PUBMED	10737800				
COMMENT	Contact: Simpson A.J.G. Laboratory of Cancer Genetics Ludwig Institute for Cancer Research Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil Tel: +55-11-2704922 Fax: +55-11-2707001 Email: asimpson@ludwig.org.br This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL				

(http://www.ludwig.org.br/scripts/gethtml2.pl?cl=cl2=RC1-HT0545-100
300-012-g05k3=2000-03-10&v=1)
Seq primer: puc18 forward
High quality sequence start: 4
High quality sequence stop: 233.
Location/Qualifiers
1. 233

FEATURES
source
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/dev_stage="Adult"
/clone_1ib="HT0545"
/note="Organ: head neck; Vector: puc18; Site 1: Sma1;
Site 2: Sma1; A mini-library was made by cloning products
derived from ORESTES PCR (U.S. Letters Patent application
No. 196,716 - Ludwig Institute for Cancer Research)
profiles into the puc18 vector. Reverse transcription of
tissue mRNA and cDNA amplification were performed under
low stringency conditions."

ORIGIN

Alignment Scores:
Pred. No.: 205 Length: 233
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x BE171313 (1-233)

OY 1 GiuguA1agInserG1yYasp 8
|||||
93 GAGGAGCCCCAGAGTGGGAGAC 116

RESULT 2
AA055968/c 301 bp mRNA linear EST 17-SEP-1996
LOCUS zf22a02.s1 Soares fetal heart NbHH19W Homo sapiens cDNA clone
DEFINITION IMAGE:377642 3', mRNA sequence.
ACCESSION AA055968
VERSION AA055968.1 GI:1548325
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 301)
Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M.,
Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M.,
Parsons, J., Rifkin, L., Rohlfing, T., Soares, M., Tan, F.,
Trevaaskis, E., Waterston, R., Williamson, A., Wohlmann, P. and
Wilson, R.

TITLE The WashU-Merck EST Project
JOURNAL Unpublished (1995)
COMMENT Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@wustl.wustl.edu
This clone is available royalty-free through LNL, contact the
IMAGE Consortium (info@image.jnl.gov) for further information.
Seg primer: -40M3 fwd. from Amerham.

FEATURES
source
1. 301
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="GDB:1285898"
/db_xref="taxon:9606"
/clone_1ib="IMAGE:377642"
/sex="unknown"
/dev_stage="19 weeks"

/lab_host="DH10B (ampicillin resistant)"
/clone_1ib="Soares fetal heart NbHH19W"
/note="Organ: heart; Vector: pRT3D (Pharmacia) with a
modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st
strand cDNA was primed with a Not I - oligo(dT) primer [5'
TGTTACCATCTGAGTGGAGCGCCGACATCTTTTCTTTTCTTTT 3']
double-stranded cDNA was size selected, ligated to Eco RI
adapters (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of a modified pRT3 vector
(Pharmacia). Library went through one round of
normalization to a cot = 5. Library constructed by
M. Fatima Bonaldo. This library was constructed from the
same fetus as the fetal lung library, Soares fetal lung
NbHH19W."

ORIGIN

Alignment Scores:
Pred. No.: 266 Length: 301
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x AA055968 (1-301)

OY 1 GiuguA1agInserG1yYasp 8
|||||
301 GAGGAGCCCCAGAGTGGGAGAC 278

RESULT 3
AA056035 327 bp mRNA linear EST 17-SEP-1996
LOCUS zf22a02.r1 Soares fetal heart NbHH19W Homo sapiens cDNA clone
DEFINITION IMAGE:377642 5', mRNA sequence.
ACCESSION AA056035
VERSION AA056035.1 GI:1548374
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 327)
Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M.,
Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M.,
Parsons, J., Rifkin, L., Rohlfing, T., Soares, M., Tan, F.,
Trevaaskis, E., Waterston, R., Williamson, A., Wohlmann, P. and
Wilson, R.

TITLE The WashU-Merck EST Project
JOURNAL Unpublished (1995)
COMMENT Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@wustl.wustl.edu
This clone is available royalty-free through LNL, contact the
IMAGE Consortium (info@image.jnl.gov) for further information.
Seg primer: -28M3 rev2 from Amerham
High quality sequence stop: 236.
Location/Qualifiers
1. 327
Location/Qualifiers

FEATURES
source
1. 327
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="GDB:1285898"
/db_xref="taxon:9606"
/clone_1ib="IMAGE:377642"
/sex="unknown"
/dev_stage="19 weeks"
/lab_host="DH10B (ampicillin resistant)"
/clone_1ib="Soares fetal heart NbHH19W"
/note="Organ: heart; Vector: pRT3D (Pharmacia) with a
modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st

strand cDNA was primed with a Not I - oligo(dT) primer [5']
 TGTACCAATCTGAAGCGAGCGCGCCGACATCTTTTCTTTTCTTTT 3'],
 double-stranded cDNA was size selected, ligated to Eco RI
 adapters (Pharmacia), digested with Not I and cloned into
 the Not I and Eco RI sites of a modified pT73 vector
 (Pharmacia). Library went through one round of
 normalization to a Cot = 5. Library constructed by
 M. Fatima Bonaldo. This library was constructed from the
 same fetus as the fetal lung library, Soares fetal lung
 NBHL19w."

ORIGIN

Alignment Scores:

Pred. No.:	290	Length:	327
Score:	41.00	Matches:	8
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	1	Gaps:	0

US-09-017-715a-2_COPY_120_127 (1-8) x AA056035 (1-327)

QY

1 GluGlulAgInSergIyGlyAsp 8
 |||||
 29 GAGGAGGCCAGAGTGGGAGAC 52

Db

RESULT 4

BM748818 333 bp mRNA linear EST 04-MAR-2002
 K-EST0023804 S9SNU601 Homo sapiens cDNA clone S9SNU601-1-C11 5',
 mRNA sequence.

LOCUS

BM748818 GI:19078436

ACCESSION

BM748818

VERSION

EST.

KEYWORDS

Homo sapiens (human)

SOURCE

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 333)

Oh,K.J., Hahn,Y., Oh,J.H., Lee,J.Y., Ahn,H.Y., Chu,M.Y., Kim,M.R.,

Oh,K.J., Cheong,J.B., Sohn,H.Y., Kim,J.M., Park,H.S., Kim,S. and

Kim,Y.S.

21C Frontier Korean EST Project 2001

Unpublished (2002)

Contact: Kim YS

Genome Research Center

Korea Research Institute of Bioscience & Biotechnology

52 Eosun-dong Yuseong-gu, Daejeon 305-333, South Korea

Tel: +82-42-860-4470

Fax: +82-42-860-4409

Email: yongsung@mail.krdb.re.kr

Plate: 1 row: C column: 11

High quality sequence stop: 333.

Location/Qualifiers

1. 333

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="S9SNU601-1-C11"

/sex="M"

/tissue_type="Ascites"

/cell_type="Epithelial"

/lab_host="STU-601"

/lab_host="TOP10"

/clone_lib="S9SNU601"

/note="Organ: Stomach; Vector: pME18-FL3; Site: 1: XhoI;

Site: 2: XhoI. The poly (A) + RNA was dephosphorylated with

bacterial alkaline phosphatase (BAP) and then decapped

with tobacco acid pyrophosphatase (TAP). The decapped

intact mRNA was ligated with DNA-RNA linker including SfiI

site by treatment of T4 RNA ligase and the first strand

cDNA was synthesized with Superscript II using SfiI

oligo-dT primer. After first strand synthesis, RNA was

degraded by NaOH treatment and cDNA was amplified by PCR
 reaction. The PCR products were digested with SfiI and
 cloned into DraIII- digested pME18-FL3 vector. The
 obtained cDNA vectors were used for transformation of
 competent cells E. coli Top10[®] by electroporation method.
 The cDNA libraries constructed by this method are
 full-length enriched cDNA library."

ORIGIN

Alignment Scores:

Pred. No.:	295	Length:	333
Score:	41.00	Matches:	8
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	4	Gaps:	0

US-09-017-715a-2_COPY_120_127 (1-8) x BM748818 (1-333)

QY

1 GluGlulAgInSergIyGlyAsp 8
 |||||
 237 GAGGAGGCCAGAGTGGGAGAC 260

Db

AI468480/c 352 bp mRNA linear EST 30-MAR-1999
 t957912.x1 NCI CGAP P-28 Homo sapiens cDNA clone IMAGE:2112934 3',
 similar to TR:015104 015104 BCSG1 PROTEIN. ; mRNA sequence.

LOCUS

AI468480

ACCESSION

AI468480

VERSION

AI468480.1 GI:4330570

KEYWORDS

EST.

SOURCE

Homo sapiens (human)

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 352)

NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

Contact: Robert Strauberg, Ph.D.

Email: sgabs-r@mail.nih.gov

Tissue Procurement: Michael J. Brownstein, M.D., Ph.D., Michael R.

Emmert-Buck, M.D., Ph.D.

cDNA Library Preparation: M. Bento Soares, Ph.D.

DNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LNL at:

www.bio.lnl.gov/bdrp/image/image.html

Insert length: 430 Std Error: 0.00

Seq primer: -40UP from Gibco

High quality sequence stop: 323.

Location/Qualifiers

1. 352

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:2112934"

/sex="male"

/dev stage="adult"

/lab_host="DH10B"

/clone_lib="NCI CGAP P-28"

/note="Organ: prostate; Vector: pT73D-Pac (Pharmacia)

with a modified polylinker; Plasmid DNA from the

normalized library NCI CGAP P-22 was prepared, and se

circles were made in vitro. Following BAP purification,

this DNA was used as tracer in a subtractive hybridization

reaction. The driver was PCR-amplified cDNAs from a pool

of 5,000 clones made from the same library (cloneids

985608-986759, 1101192-1101959, and 1217928-1220615).

Subtraction by Bento Soares and M. Fatima Bonaldo. "

ORIGIN

```

Alignment Scores:
Pred. No.:      312          Length:    352
Score:         41.00        Matches:     8
Percent Similarity:   100.00%  Conservative: 0
Best Local Similarity: 100.00%  Mismatches:  0
Query Match:       100.00%    Indels:      0
DB:                1         Gaps:           0


US-09-017-715A-2_COPY_120_127 (1-8) x AI468480 (1-352)

OY      1  gUgUAgaInserGlyVAsP 8
Db       |||||1|||||1|||||1|||||1
295 GAGAGGCCACAGACTGGGGAGAC 272

RESULT 6
LOCUS   BU729743/c              368 bp      mRNA            linear   EST 09-OCT-2002
DEFINITION
UI-E-Ckl-afj-m-04-0-Ul-sI UI-E-Ckl Homo sapiens cDNA clone
UI-E-Ckl-afj-m-04-0-Ul 3', mRNA sequence.
ACCESSION
BU729743
VERSION
BU729743.1 GI:23652933
KEYWORDS
EST.
SOURCE  Homo sapIens (human)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
1 (bases 1 to 368)
Bonaldo,M.F., Lennon,G. and Soares,M.B.
Normalization and subtraction: two approaches to facilitate gene
discovey
Genome Res. 6 (9), 791-806 (1996)
JOURNAL MEDLINE
PUBMED 9704477
COMMENT 8889548
Contact: Soares, MB
Coordinated Laboratory for Computational Genomics
University of Iowa
375 Newton Road , 4156 MEERF, Iowa City, IA 52242, USA
Tel: 319 335 8250
Fax: 319 335 9565
Email: bento-soares@uiowa.edu
Tissue Procurement: Dr. Gregg Hageman
cDNA Library preparation: Dr. M. Bento Soares, University of Iowa
cDNA library arrayed by: Dr. M. Bento Soares, University of Iowa
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
Clone Distribution: Researchers may obtain clones from Research
Genetics (www.reegen.com).
Seq primer: ML3 FORWARD
POLYA=yes
Location/Qualifiers
1..368
/oranism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="UI-E-Ckl-afj-m-04-0-Ul"
/tissue_type="Retina Foveal and Macular"
/dev_stage="adult"
/lab_host="DH10B (Life Technologies) (T1 phage resistant)"
/clone_lib="UI-E-Ckl"
/note="Organ: eye; Vector: pTR73-Pac (Pharmacia) with a
modified polylinker; Site_1: EcoR I; Site_2: Not I;
UI-E-Ckl is a normalized cDNA library containing the
following tissue(s): Retina Foveal and Macular. The
library was constructed according to Bonaldo, Lennon and
Soares, Genome Research, 6:791-806, 1996. First strand
cDNA synthesis was primed with an oligo-dT primer
containing a Not I site. Double stranded cDNA was ligated
to an EcoR I adaptor, digested with Not I, and cloned
directionally into pTR73-Pac vector. The oligonucleotide
used to prime the synthesis of first-strand cDNA contains
a library tag sequence that is located between the Not I
site and the (dT)18 tail. The sequence tag for this
library is GTCC. This library was created for the program.
```

ALIGNMENT SCORES:
 Pred. NO.: 327 Length: 368
 Score: 41.00 Matches: 8
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 5 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x BU729743 (1-368)

OY 1 GluGluAaGInserGlyGlyASP 8
 |||||
 320 GAGGAGGCCCCAGACTGGGGGAGAC 297

RESULT 7
 LOCUS AA293803/c 378 bp mRNA linear EST 12-AUG-1997
 DEFINITION IMAGE:726391 3', mRNA sequence.

ACCESSION AA293803
 VERSION AA293803.1 GI:1941726
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
 1 (bases 1 to 378)
 Hallier,L., Allen,M., Bowles,L., Dubuque,T., Geisels,G., Joest,S.,
 Kucaba,T., Lacy,M., Le,N., Lennon,G., Maira,M., Martin,J.,
 Moore,B., Schellenberg,K., Seepce,M., Tan,F., Theising,B.,
 White,Y., Wyile,T., Waterston,R. and Wilson,R.
 WashU-Merck EST Project 1997
 Unpublished (1997)
 CONTACT: Wilson RK
 Washington University School of Medicine
 444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: est@watson.wustl.edu
 This clone is available royalty-free through LINT ; contact the
 IMAGE Consortium (info@image.llnl.gov) for further information.
 Insert length: 484 Spd Error: 0.00
 Seq primer: -41m13 fwd, RT from Amersham.

LOCATION/QUALIFIERS
 1..378
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="GDB:5938504"
 /db_xref="taxon:9606"
 /clone="IMAGE:726391"
 /sex="Female"
 /tissue_type="ovarian tumor"
 /lab_host="DH10B (ampicillin resistant)"
 /clone_lib="Soares ovary tumor NBHOT"

/note="Organ: ovary; Vector: pT7T3D (Pharmacia) with a
 modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st
 strand cDNA was primed with a Not I - oligo(dT) primer [5'
 TGTTACCAACTGTAAGTGGGCGCGCGCGCTTTTCTTTTCTTTTCTT 3'],
 double-stranded cDNA was size selected, ligated to Eco RI
 adapters (Pharmacia), digested with Not I and cloned into
 the Not I and Eco RI sites of a modified pT7T3 vector
 (Pharmacia). Library constructed by Bento Soares and
 M.Fatima Bonaído."

Pred. No.: 336 Length: 378
 Score: 41.00 Matches: 8
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 1 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x AA293803 (1-378)

QY 1 GluGlu1aGlnSerg1yG1yASP 8
 DB 302 GAGGAGGCCAGACTGGGAGAGAC 279

RESULT 8
 AA722407 387 bp mRNA linear EST 02-JAN-1998
 LOCUS 2983h10.81 Soares fetal heart NBH19W Homo sapiens cDNA clone
 DEFINITION IMAGE:400003 3', mRNA sequence.

ACCESSION AA722407
 VERSION AA722407
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens

REFERENCE
 AUTHORS Hillier, L., Allen, M., Bowles, L., Dubuque, T., Getse, G., Joet, S., Kitzman, D., Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin, J., Moore, B., Scheinberg, K., Stepien, M., Tan, F., Theising, B., White, Y., Wylie, T., Waterston, R. and Wilson, R.
 TITLE Washu-Merck EST Project
 JOURNAL Unpublished (1997)
 COMMENT Contact: Wilson RK
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: est@watson.wustl.edu

FEATURES
 source
 This clone is available royalty-free through LNL; contact the IMAGE Consortium (info@image.lnl.gov) for further information.
 Seq primer: -40m13 fwd. ET from Amersham
 High quality sequence stop: 384.
 Location/Qualifiers
 1..387
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="GDB:1307826"
 /db_xref="taxon:9606"
 /clone="IMAGE:400003"
 /sex="unknown"
 /dev_stage="19 weeks"
 /lab_host="DH10B (ampicillin resistant)"
 /clone_1lb="Soares fetal heart NBH19W"
 /note="Organ: heart; Vector: pT73D (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5' TGTTACCAATCTGAAGTGGAGCGCGGCGATCTTTTCTTTTCTTTT 3'], double-stranded cDNA was size selected, ligated to Eco RI adapters (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of a modified pT73 vector (Pharmacia). Library went through one round of normalization to a Cot = 5. Library constructed by M.Fatima Bernaldo. This library was constructed from the same fetus as the fetal lung library, Soares fetal lung NBH19W."

ORIGIN

Alignment Scores:
 Pred. No.: 344 Length: 387
 Score: 41.00 Matches: 8
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0

DB: 1 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x AA722407 (1-387)

QY 1 GluGlu1aGlnSerg1yG1yASP 8
 DB 297 GAGGAGGCCAGACTGGGAGAGAC 274

RESULT 9
 AA394097 393 bp mRNA linear EST 12-AUG-1997
 LOCUS z156h04.r1 Soares ovary tumor NBH07 Homo sapiens cDNA clone
 DEFINITION IMAGE:726391 5', similar to TR:G971580 G971580 SENSORY NEURON SYNUCLEIN, mRNA sequence.

ACCESSION AA394097
 VERSION AA394097
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens

REFERENCE
 AUTHORS Hillier, L., Allen, M., Bowles, L., Dubuque, T., Getse, G., Joet, S., Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin, J., Moore, B., Scheinberg, K., Stepien, M., Tan, F., Theising, B., White, Y., Wylie, T., Waterston, R. and Wilson, R.
 TITLE Washu-Merck EST Project 1997
 JOURNAL Unpublished (1997)
 COMMENT Contact: Wilson RK
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: est@watson.wustl.edu

FEATURES
 source
 This clone is available royalty-free through LNL; contact the IMAGE Consortium (info@image.lnl.gov) for further information.
 Insert length: 484 Std Error: 0.00
 Seq primer: -28m13 rev2 ET from Amersham.
 Location/Qualifiers
 1..393
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="GDB:5938504"
 /db_xref="taxon:9606"
 /clone="IMAGE:726391"
 /sex="female"
 /tissue_type="ovarian tumor"
 /lab_host="DH10B (ampicillin resistant)"
 /clone_1lb="Soares ovary tumor NBH07"
 /note="Organ: ovary; Vector: pT73D (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5' TGTTACCAATCTGAAGTGGAGCGCGGCGGATCTTTTCTTTTCTTTT 3'], double-stranded cDNA was size selected, ligated to Eco RI adapters (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of a modified pT73 vector (Pharmacia). Library constructed by Bento Soares and M.Fatima Bernaldo."

ORIGIN

Alignment Scores:
 Pred. No.: 349 Length: 393
 Score: 41.00 Matches: 8
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 1 Gaps: 0
 US-09-017-715A-2_COPY_120_127 (1-8) x AA394097 (1-393)
 QY 1 GluGlu1aGlnSerg1yG1yASP 8
 DB 111 GAGGAGGCCAGACTGGGAGAGAC 134

RESULT 10
CF529590/c 404 bp mRNA linear EST 11-SEP-2003
LOCUS UI-1-BC1P-asy-b-08-0-UI.s1 NCI CGAP_P13 Homo sapiens cDNA clone
DEFINITION UI-1-BC1P-asy-b-08-0-UI 3', mRNA sequence.
ACCESSION CF529590
VERSION CF529590.1 GI:34579285
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homindae; Homo.
REFERENCE NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncigap>.
1 (bases 1 to 404)
AUTHORS National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
TITLE Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cga@bbs-rcmail.nih.gov
Tissue Procurement: Dr. Steven Brown
cDNA Library Preparation: Dr. M. Bento Soares, University of Iowa
cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
Clone Distribution: Clone distribution information can be obtained
from Dr. M. Bento Soares, bento-soares@uiowa.edu
Seq primer: M13 FORWARD
POLYA=Yes.

FEATURES
source
1..404
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="UI-1-BC1P-asy-b-08-0-UI"
/tissue_type="Placenta"
/dev_stage="8-9 weeks"
/lab_host="DH10B (Life Technologies)"
/clone_1ib="NCI CGAP_P13"
/note="Organ: Placenta; Vector: pT73-Pac (Pharmacia) with
a modified polylinker; Site_1: EcoR I; Site_2: Not I;
NCI CGAP_P13 is a subcloned cDNA library constructed
according to Bonaldo, Lennon and Soares, Genome Research,
6:791-806, 1996. First strand cDNA synthesis was primed
with an oligo-dT primer containing a Not I site. Double
stranded cDNA was ligated to an EcoR I adaptor, digested
with Not I, and cloned directionally into pT73-Pac
vector. The oligonucleotide used to prime the synthesis of
first-strand cDNA contains a library tag sequence that is
located between the Not I site and the (dT)18 tail. The
sequence tags for this library are GA, AGGA. For
additional information, contact: Bento Soares,
bento-soares@uiowa.edu
TAG TISSUE=Placenta human 8 week
TAG LIB=UI-1-BC1P
TAG_SEQ=GA"

ORIGIN
Alignment Scores:
Pred. No.: 359 Length: 404
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 7 Gaps: 0
US-09-017-715A-2_COPY_120_127 (1-8) x CF529590 (1-404)
QY 1 GUGUUAAGlnSerglyasp 8
DB 321 GAGGAGGCCAGAGTGGGAGAC 298
RESULT 11
AV683707
AV683707

LOCUS AV683707 407 bp mRNA linear EST 16-JAN-2002
DEFINITION AV683707 GKC Homo sapiens cDNA clone GKCEPB06 5', mRNA sequence.
ACCESSION AV683707
VERSION AV683707.1 GI:10285570
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homindae; Homo.
REFERENCE 1 (bases 1 to 407)
AUTHORS Xu, X., Huang, J., Xu, Z., Qian, B., Zhu, Z., Yan, Q., Cai, T., Zhang, X.,
Xiao, H., Qu, J., Liu, F., Huang, Q., Cheng, Z., Li, N., Du, J., Hu, W.,
Shen, K., Lu, G., Fu, G., Zhong, M., Xu, S., Gu, W., Huang, W., Zhao, X.,
Hu, G., Gu, J., Chen, Z., and Han, Z.
TITLE Insight into hepatocellular carcinogenesis at transcriptome level
by comparing gene expression profiles of hepatocellular carcinoma
with those of corresponding noncancerous liver
Proc. Natl. Acad. Sci. U.S.A. 98 (26), 15085-15094 (2001)
JOURNAL 11752456
MEDLINE 21625106
PUBMED
COMMENT Contact: Zeguang Han
Chinese National Human Genome Center at Shanghai
351 Guo Shoujing Road, Zhangjiang Hi-Tech Park, Pudong, Shanghai
201203, P. R. China
Tel: 86-21-50801919 (ex. 45)
Fax: 86-21-50801922
Email: hanzg@chgc.sh.cn
This clone is available at CHGC in Shanghai.

FEATURES
source
1..407
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="GKCEPB06"
/tissue_type="hepatocellular carcinoma"
/dev_stage="Adult"
/lab_host="SOLR"
/clone_1ib="GKC"
/note="Vector: pBluescript sk(-); Site_1: EcoRI; Site_2:
XhoI"

ORIGIN
Alignment Scores:
Pred. No.: 362 Length: 407
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 1 Gaps: 0
US-09-017-715A-2_COPY_120_127 (1-8) x AV683707 (1-407)
QY 1 GUGUUAAGlnSerglyasp 8
DB 169 GAGGAGGCCAGAGTGGGAGAC 192
RESULT 12
AA946606/c 408 bp mRNA linear EST 23-JUL-1998
LOCUS AA946606
DEFINITION cq38c12.s1 NCI CGAP Kids Homo sapiens cDNA clone IMAGE:1588630 3'
similar to TR:O15104 O15104 BCSG1 PROTEIN. ;, mRNA sequence.
ACCESSION AA946606
VERSION AA946606.1 GI:3110001
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homindae; Homo.
REFERENCE 1 (bases 1 to 408)
AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncigap>.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)

COMMENT

Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: M. Bento Soares, Ph.D.
DNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bdip/image/image.html
Insert Length: 812 Std Error: 0.00
Seq primer: -40m13 fwd. RT from Amersham
High quality sequence stop: 359.

FEATURES

source

```
1..408
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:1588630"
/tissue_type="2 pooled tumors (clear cell type)"
/lab_host="DH10B"
/clone_id="NCI CGAP Kid5"
/note="Organ: kidney; Vector: pTV73D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5' ACTGGAAGATCGCGCGCATTTTCTTTTCTTTT 3'], double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pTV73 vector. Library went through one round of normalization. Library constructed by Bento Soares and M. Fatima Bonaldo. "
```

ORIGIN

Alignment Scores:

Pred. No.:	363	Length:	408
Score:	41.00	Matches:	8
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	1	Gaps:	0

US-09-017-715A-2_COPY_120_127 (1-8) x AA946606 (1-408)

QY 1 GUGUUAAGUInsergyGlyAap 8

Db 309 GAGGAGGCCAGAGTGGGAGAC 286

RESULT 13

A1139933/c

LOCUS

DEFINITION

A1139933 412 bp mRNA linear EST 05-OCT-1998
g668B04.X1 Soares_fetal_heart_NBH119W Homo sapiens cDNA clone
IMAGE:1691887 3 similar to TR:015104 015104 BCSG1 PROTEIN. ; mRNA
sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 584 Std Error: 0.00
Seq primer: -40m13 fwd. RT from Amersham.
Location/Qualifiers
1..412

ORIGIN

```
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:1691887"
/sex="unknown"
/dev_stage="19 weeks"
/lab_host="DH10B (ampicillin resistant)"
/clone_id="Soares_fetal_heart_NBH119W"
/note="Organ: heart; Vector: pTV73D (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5' TGTTCACCATCGAAGCGAGCGCGCATTTTCTTTTCTTTT 3'], double-stranded cDNA was size selected, ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of a modified pTV73 vector (Pharmacia). Library went through one round of normalization to a Cot = 5. Library constructed by M. Fatima Bonaldo. This library was constructed from the same fetus as the fetal lung library, Soares fetal lung NBH119W."
```

Alignment Scores:

Pred. No.:	367	Length:	412
Score:	41.00	Matches:	8
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	1	Gaps:	0

US-09-017-715A-2_COPY_120_127 (1-8) x A1139933 (1-412)

QY 1 GUGUUAAGUInsergyGlyAap 8

Db 303 GAGGAGGCCAGAGTGGGAGAC 280

RESULT 14

AV703171

LOCUS

DEFINITION

AV703171 421 bp mRNA linear EST 09-OCT-2000
AV703171 ADB Homo sapiens cDNA clone ADBBCF03 5', mRNA sequence.
AV703171.1 GI:10720500

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Unpublished (2000)
Contact: Zengqiang Han
Chinese National Human Genome Center at Shanghai
351 Guo Shoujing Road, Zhangjiang Hi-Tech Park, Pudong, Shanghai
201203, P. R. China
Tel: 86-21-50801919 (ex.45)
Fax: 86-21-50801922
Email: hanzg@chgc.sh.cn
This clone is available at CHGC in Shanghai.
Location/Qualifiers
1..421

```
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="ADBBCF03"
/tissue_type="Adrenal gland"
/dev_stage="Adult"
/lab_host="SOLR"
/clone_id="ADB"
/note="Vector: pBluescript sk(-); Site 1: EcoRI; Site 2:
```

ORIGIN XhoI"

Alignment Scores:

Pred. No.:	375	Length:	421
Score:	41.00	Matches:	8
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	1	Gaps:	0

US-09-017-715A-2_COPY_120_127 (1-8) x AV703171 (1-421)

Qy 1 GIJGUAlaGInserGlyGlyASP 8

DB 374 GAGGAGCCCGACAGTGGGGAGAC 397

RESULT 15

BX090816

428 bp mRNA linear EST 23-JAN-2003

LOCUS BX090816 Soares ovary tumor NBHOT Homo sapiens cDNA clone

DEFINITION IMAGP998P081781 ; IMAGE:726391, mRNA sequence.

ACCESSION BX090816

VERSION BX090816.1 GI:27822248

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 428)

AUTHORS Ebert, L., Heil, O., Hennig, S., Neubert, P., Partsch, E., Peters, M.,

TITLE Human Unigeneset - RZPD3

JOURNAL Unpublished (2003)

COMMENT Contact: Ina Rolfs

RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH

Im Neuenheimer Feld 580, D-69120 Heidelberg, Germany

RZPD; IMAGP998P081781.

RZPDLIB; I.M.A.G.E. cDNA Clone Collection:

http://www.rzpd.de/Cloncards/cgi-

bin/showLib.pl.cgi?response?libno=972 Contact: Ina Rolfs

RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH

Heubnerweg 6, D-14059 Berlin, Germany

Tel: +49 30 32639 101

Fax: +49 30 32639 111

www.rzpd.de

This clone is available royalty-free from RZPD;

contact RZPD (clone@rzpd.de) for further information. Seq primer:

M13r, Primer sequence: TTTCAACACGAGAAACAGCTATGAC.

Location/Qualifiers

1. 428

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGP998P081781 ; IMAGE:726391"

/sex="Female"

/tissue_type="ovarian tumor"

/lab_host="DH10B (ampicillin resistant)"

/clone_lib="Soares ovary tumor NBHOT"

/note="Organ: ovary; Vector: p773D (Pharmacia) with a

modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st

strand cDNA was primed with a Not I - oligo(dT) primer (5'

TGTTACCATCTGAGTGGAGCGCGGTTTTTTTTTTTTTTT 3').

double-stranded cDNA was size selected, ligated to Eco RI

adapters (Pharmacia), digested with Not I and cloned into

the Not I and Eco RI sites of a modified p773 vector

(Pharmacia). Library constructed by Bento Soares and

M.Fatima Bonaldo."

Score:	41.00	Matches:	8
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	5	Gaps:	0

US-09-017-715A-2_COPY_120_127 (1-8) x BX090816 (1-428)

Qy 1 GIJGUAlaGInserGlyGlyASP 8

DB 111 GAGGAGCCCGACAGTGGGGAGAC 134

Search completed: May 4, 2005, 13:46:36
 Job time : 364.973 secs

ORIGIN

Alignment Scores:

Pred. No.: 381

Length: 428

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - nucleic search, using frame_plus_p2n model

Run on: May 4, 2005, 09:07:56 ; Search time 636.953 Seconds

(without alignments)
836.639 Million cell updates/sec

Title: US-09-017-715A-2_COPY_94_107
Sequence: 1 VVRKEDLRSPAPQ 14

Scoring table:
BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 45 summaries

Command line parameters:

-MODEL=frame+g2n.model -DEV=xlh
-O=/cg92.1/USPTO.spool.h/US09017715/runat_04052005_100744_25619/app_query.fasta_1.661
-DB=EST -OPMT=fastrap -SUFFIX=rcs -MINMATCH=0.1 -LOOPCL=0 -LOOPEXT=0
-UNITS=bits -START=1 -END=1 -MATRIX=BLOSUM62 -TRANS=human40.cdi -LIST=45
-LOCALIGN=200 -THR SCORE=pct -THR MAX=100 -THR MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTPM=pcr -NOR=exc -HEAPSIZE=500 -MINLEN=0 -MAXLEN=200000000
-USER=US09017715_@CGN_1_1_5334_@runat_04052005_100744_25619 -NCPU=6 -ICPU=3
-NO_MMAP -LARGEQUERY -NEG_SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG
-DEV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

EST:
1: gb_esc1.*
2: gb_esc2.*
3: gb_hlc.*
4: gb_esc3.*
5: gb_esc4.*
6: gb_esc5.*
7: gb_esc6.*
8: gb_gsa2.*
9: gb_gsa2.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	70	100.0	161	2 BE774132	MR1-UM000
2	70	100.0	283	4 BM693306	UI-E-CK1
3	70	100.0	333	4 BM748818	K-EST0023
4	70	100.0	369	1 AV708398	AV708398
5	70	100.0	387	1 AA722407	Z963h10.8
6	70	100.0	393	1 AA394097	z56h04.x
7	70	100.0	407	1 AA683707	AV683707
8	70	100.0	412	1 A1139933	qae8b04.x
9	70	100.0	421	1 AV703171	AV703171

10	70	100.0	428	5 BX090816	BX090816
C 11	70	100.0	438	1 A1684600	A1684600
C 12	70	100.0	442	1 AA804675	0144b01.8
C 13	70	100.0	451	4 BG826435	602750062
C 14	70	100.0	462	5 BX474511	BKF2p686E
C 15	70	100.0	467	6 CB107161	K-EST0145
C 16	70	100.0	472	4 BM704200	UI-E-CK1
C 17	70	100.0	474	4 BM706956	UI-E-CK1
C 18	70	100.0	480	4 BM68990	UI-E-CQ0
C 19	70	100.0	489	5 BP201709	BP201709
C 20	70	100.0	504	5 BU728272	UI-E-CQ0
C 21	70	100.0	509	2 BE299889	600944691
C 22	70	100.0	510	2 BE298825	601119383
C 23	70	100.0	519	1 A1016464	0178h06.8
C 24	70	100.0	538	4 BM695726	UI-E-CQ1
C 25	70	100.0	544	4 BM665098	UI-E-CQ1
C 26	70	100.0	555	5 BX474500	DKE2p686D
C 27	70	100.0	555	7 CV028548	7090_Fu11
C 28	70	100.0	558	4 B1548891	603189023
C 29	70	100.0	568	4 BG708703	602674249
C 30	70	100.0	578	5 BP212912	BP212912
C 31	70	100.0	582	5 BP197662	BP197662
C 32	70	100.0	583	5 BP200612	BP200612
C 33	70	100.0	584	1 AV708933	AV708933
C 34	70	100.0	584	5 BP201686	BP201686
C 35	70	100.0	588	4 B1755243	603023269
C 36	70	100.0	592	6 CA443299	UI-H-DH1
C 37	70	100.0	593	6 CA443166	UI-H-DH1
C 38	70	100.0	614	5 BU730570	UI-E-C11
C 39	70	100.0	617	1 AA633976	ac33e03.8
C 40	70	100.0	617	4 BG707764	602671103
C 41	70	100.0	623	5 BP381244	BP381244
C 42	70	100.0	641	1 AL712443	DKE2p686B
C 43	70	100.0	653	4 B1757131	603030894
C 44	70	100.0	659	7 CN410061	170006001
C 45	70	100.0	706	4 BG328738	602427017

ALIGNMENTS

RESULT 1	BE774132	161 bp	MRNA	linear	EST 20-SEP-2000
LOCUS	BE774132				
DEFINITION	MR1-UM0009-220500-015-g07_1 UM0009 Homo sapiens cDNA, mRNA				
ACCESSION	BE774132				
VERSION	BE774132.1	GI:10227839			
KEYWORDS	EST.				
SOURCE	Homo sapiens (human)				
ORGANISM	Homo sapiens				
REFERENCE	Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R., Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F., Goldman,G.H., Carvalho,A.P., Matsunuma,A., Baia,G.S., Simpson,D.H., Brunstein,A., deOliveira,P.S., Bucher,P., Jorgensen,C.V., O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and Simpson,A.J.				
AUTHORS	Shotgun sequencing of the human transcriptome with ORF expressed				
TITLE	sequence tags				
JOURNAL	Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)				
MEDLINE	20202663				
PUBMED	10737800				
COMMENT	Contact: Simpson A.J.G. Laboratory of Cancer Genetics Ludwig Institute for Cancer Research Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil Tel: +55-11-2704922 Fax: +55-11-2707001 Email: asimpson@ludwig.org.br This sequence was derived from the FAPESP/LICR Human Genome				

Project. This entry can be seen in the following URL
(<http://www.ludwig.org.br/scripts/gethtml2.pl?cl=6c2-MR1-UM0009-220>)
500-015-g01_let3=2000-05-22&cl=1)
Seq primer: puc 18 forward

High quality sequence stop: 161.
Location/Qualifiers

1. 161

/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/dev_stage="Adult"
/clone_lib="UM0009"
/note="Organ: uterus; Vector: puc18; Site 1: SmaI; Site 2: SmaI; A mini-library was made by cloning products derived from ORESTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research) profiles into the pUC 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."

ORIGIN

Alignment Scores:

Pred. No.:	0.0174	Length:	161
Score:	70.00	Matches:	14
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	2	Gaps:	0

US-09-017-715a-2_COPY_94_107 (1-14) x BE74132 (1-161)

Qy 1 ValValArgLyGluAspLeuArgProSerAlaProGlnGln 14

Db 69 GTGGTCCGACAGAGACTTGAGCCATCTGCCCCCAACAG 110

RESULT 2

LOCUS

BM693306 283 bp mRNA linear EST 28-FEB-2002

DEFINITION UI-E-CK1-afm-1-14-0-UI r1 UI-E-CK1 Homo sapiens cDNA clone

ACCESSION BM693306

VERSION BM693306.1 GI:19006564

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

TITLE 1 (bases 1 to 283)

Normalisation and subtraction: two approaches to facilitate gene

discovery

JOURNAL Genome Res. 6 (9), 791-806 (1996)

MEDLINE 97044477

COMMENT 8889548

CONTACT: Soares, MB

Coordinated Laboratory for Computational Genomics

University of Iowa

375 Newton Road, 4156 MEHRF, Iowa City, IA 52242, USA

Tel: 319 335 8250

Fax: 319 335 9565

Email: bento-soares@iowa.edu

Tissue Procurement: Dr. Gregg Hageman

cDNA library preparation: Dr. M. Bento Soares, University of Iowa

cDNA library Arrayed by: Dr. M. Bento Soares, University of Iowa

DNA Sequencing by: Dr. M. Bento Soares, University of Iowa

Clone Distribution: Researchers may obtain clones from Research

Genetics (www.resgen.com).

Seq primer: M13 Reverse.

Location/Qualifiers

1. 283

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/cell_type="Epithelial"

/clone="UI-E-CK1-afm-1-14-0-UI"
/tissue_type="Retina Foveal and Macular"
/dev_stage="adult"
/lab_host="DH10B (Life Technologies) (T1 phage resistant)"
/clone_lib="UI-E-CK1"
/note="Organ: eye; Vector: pT73-Pac (Pharmacia) with a modified polylinker; Site 1: EcoR I; Site 2: Not I; UI-E-CK1 is a normalized cDNA library containing the following tissue(s): Retina Foveal and Macular. The library was constructed according to Donald, Lennon and Soares, Genome Research, 6:791-806, 1996. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was ligated to an EcoR I adaptor, digested with Not I, and cloned directionally into pT73-Pac vector. The oligonucleotide used to prime the synthesis of first-strand cDNA contains a library tag sequence that is located between the Not I site and the (dT)18 tail. The sequence tag for this library is GTCC. This library was created for the program, Gene Discovery in the Visual System, supported by National Eye Institute (NEI)."

ORIGIN

Alignment Scores:

Pred. No.:	0.0316	Length:	283
Score:	70.00	Matches:	14
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	4	Gaps:	0

US-09-017-715a-2_COPY_94_107 (1-14) x BM693306 (1-283)

Qy 1 ValValArgLyGluAspLeuArgProSerAlaProGlnGln 14

Db 229 GTGGTCCGACAGAGACTTGAGCCATCTGCCCCCAACAG 270

RESULT 3

LOCUS

BM748818 333 bp mRNA linear EST 04-MAR-2002

DEFINITION K-EST0023804 S9SNU601 Homo sapiens cDNA clone S9SNU601-1-C11 5',

ACCESSION BM748818

VERSION BM748818.1 GI:19078436

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

TITLE 1 (bases 1 to 333)

Normalisation and subtraction: two approaches to facilitate gene

discovery

JOURNAL Oh K.J., Cheong, J.E., Sohn, H.Y., Kim, J.M., Park, H.S., Kim, S. and

Kim, Y.S.

21C Frontier Korean EST Project 2001

Unpublished (2002)

CONTACT: Kim YS

Genome Research Center

Korea Research Institute of Bioscience & Biotechnology

52 Boeun-dong Yuseong-gu, Daejeon 305-333, South Korea

Tel: +82-42-860-4470

Fax: +82-42-860-4409

Email: yongsung@mail.kribb.re.kr

Plate: 1 row: C column: 11

High quality sequence stop: 333.

Location/Qualifiers

1. 333

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="S9SNU601-1-C11"

/sex="M"

/tissue_type="Ascites"

/cell_type="Epithelial"

```

/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/_clone="ADCMAB05"
/_issue_type="Adrenal gland"
/_dev_stage="Adult"
/_lab_host="SOLR"
/_clone_lid="ADC"
/_note="Vector: pBluescript sk(-), Site_1: EcoRI, Site_2
XhoI"

```

Alignment Scores:
Pred. No.:
Score:
Percent Similarity:

0.044	Length:	387
70.00	Matches:	14
100.00%	Conservative:	0

Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 1 Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x AA722407 (1-387)

Qy 1 ValValaJrlyGjUaSpLeuArProSeRaIaProGInGln 14
|||||
375 GTGGTCCGCAAGAGACTTGAGCCATCTGCCCCCAACAG 334

RESULT 6
AA394097 393 bp mRNA linear EST 12-AUG-1997
LOCUS z55h04.x1 Soares ovary tumor NbH09 Homo sapiens cDNA clone
DEFINITION IMAGE:726391.5, similar to TR:G971580 G971580 SENSORY NEURON
SYNCDLEIN.?, mRNA sequence.

ACCESSION AA394097
VERSION AA394097.1 GI:2047067
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 393)
AUTHORS Hillier, L., Allen, M., Bowles, L., Dubuque, T., Geisel, G., Jost, S.,
Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin, J.,
Moore, B., Schellenberg, K., Steptoe, M., Tan, F., Theisling, B.,
White, Y., Wylie, T., Waterston, R. and Wilson, R.
WashU-Merck EST Project 1997
Unpublished (1997)

TITLE
JOURNAL
COMMENT Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810

FEATURES
source
1.393
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="GDB:5938504"
/db_xref="taxon:9606"
/clone="IMAGE:726391"
/sex="Female"
/tissue_type="ovarian tumor"
/lab_host="DH10B (ampicillin resistant)"
/clone_lib="Soares ovary tumor NbH09"
/note="Organ: ovary; Vector: pRT73D (Pharmacia) with a
modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st
strand cDNA was primed with a Not I - oligo (dT) primer [5'
TGTTACCAATCTGAAGTGGAGCGCGCTTTTCTTTTCTTTT 3'],
double-stranded cDNA was size selected, ligated to Eco RI
adapters (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of a modified pRT73 vector
(Pharmacia). Library constructed by Bento Soares and
M.Fatima Bonaldo."

ORIGIN

Alignment Scores:

Pred. No.: 0 0.447 Length: 393
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 1 Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x AA394097 (1-393)

Qy 1 ValValaJrlyGjUaSpLeuArProSeRaIaProGInGln 14

DB 33 GTGGTCCGCAAGAGACTTGAGCCATCTGCCCCCAACAG 74
|||||

RESULT 7

AV683707 407 bp mRNA linear EST 16-JAN-2002
LOCUS AV683707 GKC Homo sapiens cDNA clone GKCPB06 5', mRNA sequence.

DEFINITION AV683707
ACCESSION AV683707
VERSION AV683707.1 GI:10285570
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 407)
AUTHORS Xu, X., Huang, J., Xu, Z., Qian, B., Zhu, Z., Yan, Q., Cai, T., Zhang, X.,
Xiao, H., Qu, J., Liu, F., Huang, Q., Cheng, Z., Li, N., Du, J., Hu, W.,
Shen, K., Lu, G., Fu, G., Zhong, M., Xu, S., Gu, W., Huang, W., Zhao, X.,
Hu, G., Gu, J., Chen, Z., and Han, Z.

TITLE
JOURNAL
MEDLINE
PUBMED
COMMENT Contact: Zeguang Han
Chinese National Human Genome Center at Shanghai
351 Guo Shoujing Road, Zhangjiang Hi-Tech Park, Pudong, Shanghai
201203, P. R. China
Tel: 86-21-50801919 (ex. 45)
Fax: 86-21-50801922
Email: hanzg@chgc.sh.cn
This clone is available at CHGC in Shanghai.

FEATURES
source
1.407
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="GKCPB06"
/tissue_type="hepatocellular carcinoma"
/dev_stage="Adult"
/lab_host="SOLR"
/clone_lib="GKC"
/note="Vector: pBluescript sk(-); Site_1: EcoRI; Site_2:
XhoI"

ORIGIN

Alignment Scores:

Pred. No.: 0.0464 Length: 407
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 1 Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x AV683707 (1-407)

Qy 1 ValValaJrlyGjUaSpLeuArProSeRaIaProGInGln 14
|||||
94 GTGGTCCGCAAGAGACTTGAGCCATCTGCCCCCAACAG 135

RESULT 8

AI139933/c 412 bp mRNA linear EST 05-OCT-1998
LOCUS qae6b04.x1 Soares fetal heart NbH119W Homo sapiens cDNA clone
DEFINITION IMAGE:1691887 3', similar to TR:O15104 O15104 BCSG1 PROTEIN.?, mRNA
sequence.

ACCESSION AI139933
VERSION AI139933.1 GI:3647390
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE AUTHORS TITLE	DOI
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo. 1 (baes 1 to 412) NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap .	10.1002/ajpa.20001
National Cancer Institute, Cancer Genome Anatomy Project (CGAP)	10.1002/ajpa.20001

Alignment Scores:

Pred. No.:	0.047	length:	4122
Score:	70.00	Matches:	14
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0

US-09-017-715A-2_COPY_94_107 (1-14) X A1139933 (1-412)

QY 1 ValValArglyGluAspLeuArgProSerAlaProGlnGln 14
DB 381 GTGTGCGCAAGAGACTTGAGGCCATCTGCCCCCAACAG 340

RESULT 9
AV703171

REFERENCE
AUTHORS
1 (bases 1 to 421)
Peng, Y., Song, H., Huang, Q., Huang, C., Gu, Y., Yang, Y., Gao, G.,

TITLE
JOURNAL
COMMENT

FEATURES

- source
- 1. 421
- Location/Qualifiers
- This clone is available at CHGC in Shanghai
- Email: hanzg@chgc.sh.cn
- Fax: 86-21-50801922

ORIGIN

Alignment Scores:

Pred. No.:	0.0481	Length:	422
Score:	70.00	Matches:	14
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	1	Gaps:	0

US-09-017-715A-2_COPY_94_107 (1-14) X AV703171 (1-421)

QY 1 ValValArgLysGlaspleuArgProSerAlaProGlnGln 14
| | | | | | | | | | | | | | | | | |
Db 296 GTGTGCGCAAGAGAATTGAGGCCATCTGCCCCCACAAG 33

LOCUS	428 bp	mRNA	linear	EST 23-JAN-2003
DEFINITION	BX030816			
	BX030816	Soares	ovary tumor NBOHT Homo sapiens	CDNA clone
	IMAGE998EP081781		IMAGE:726391	mRNA sequence.

REFERENCE	AUTHORS	TITLE	JOURNAL	COMMENT
1 (bases 1 to 428)	Eberl, L., Heil, O., Hennig, S., Neubert, P., Partsch, E., Peters, M., Radehof, U., Schneider, D. and Korn, B.	Human Unigeneset - RZPD3	Unpublished (2003)	Contact: Ina Rolfs

FEATURES
SOURCE

```

/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGP938P081781 ; IMAGE:725391"
/sex="Female"
/tissue_type="ovarian tumor"
/tab_host="DH10B (ampicillin resistant)

```

/clone.lib="Soares ovary tumor NBH07"
 /note="Organ: ovary; Vector: pRT73D (Pharmacia) with a
 modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st
 strand cDNA was primed with a Not I - oligo (dT) primer (5'
 TGTTCACATCTGAGAGGAGCGCGGGGTTTTTTTTTTTTTTT 3'),
 double-stranded cDNA was size selected, ligated to Eco RI
 adapters (Pharmacia), digested with Not I and cloned into
 the Not I and Eco RI sites of a modified pRT73 vector
 (Pharmacia). Library constructed by Bento Soares and
 M. Fatima Bonaldo."

ORIGIN

Alignment Scores:

Pred. No.:	0.0489	Length:	428
Score:	70.00	Matches:	14
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	5	Gaps:	0

US-09-017-715A-2_COPY_94_107 (1-14) x BK090816 (1-428)

Qy 1 ValValArgLySGluAspLeuArgProSerAlaProGlnGln 14
 |||||
 Db 33 GTGGTGGCGCAGAGAGACTTGAGGCGCATCTGCCCCCAACAG 74

RESULT 11

AI684600 438 bp mRNA linear EST 07-MAR-2000
 LOCUS wa84d12.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone
 DEFINITION IMAGE:2302871 3' similar to TR:015104 O15104 BCSG1 PROTEIN. [1] ;
 mRNA sequence.

ACCESSION AI684600.1 GI:4895894
 VERSION EST.
 KEYWORDS Homo sapiens (human)
 SOURCE Homo sapiens

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 438)

NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-remail.nih.gov

This clone is available royalty-free through LML; contact the

IMAGE Consortium (info@image.llnl.gov) for further information.

Insert Length: 505 Std Error: 0.00

Seg primer: -40UP from Gibco.

Location/Qualifiers

FEATURES

source

1..438
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:2302871"
 /lab_host="DH10B"

/clone.lib="Soares_NFL_T_GBC_S1"

/note="Organ: pooled; Vector: pRT73D-Pac (Pharmacia) with
 a modified polylinker; Site_1: Not I; Site_2: Eco RI;
 Equal amounts of plasmid DNA from three normalized

libraries (fetal lung NBH19W, testis NHT, and B-cell

NCI CGAP GCBI) were mixed, and as circles were made in

vitro. Following HAP purification, this DNA was used as

tracer in a subtractive hybridization reaction. The driver

was PCR-amplified cDNAs from pools of 5,000 clones made

from the same 3 libraries. The pools consisted of

I.M.A.G.E. clones 297480-302087, 682632-687239,

726408-728711, and 728096-731399. Subtraction by Bento

Soares and M. Fatima Bonaldo."

ORIGIN

Alignment Scores:

Pred. No.:	0.0502	Length:	438
Score:	70.00	Matches:	14
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	1	Gaps:	0

US-09-017-715A-2_COPY_94_107 (1-14) x AI684600 (1-438)

Qy 1 ValValArgLySGluAspLeuArgProSerAlaProGlnGln 14
 |||||
 Db 378 GTGGTGGCGCAGAGAGACTTGAGGCGCATCTGCCCCCAACAG 337

RESULT 12

AA804675 442 bp mRNA linear EST 19-FEB-1998
 LOCUS oF44b01.s1 NCI_CGAP CNS1 Homo sapiens cDNA clone IMAGE:1427017 3'
 DEFINITION similar to SW:SYUN_RAT Q63544 SENSORY NEURON SYNUCLEIN. ; mRNA
 sequence.

ACCESSION AA804675.1 GI:2876076

VERSION EST.

KEYWORDS Homo sapiens (human)

SOURCE Homo sapiens

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 442)

NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-remail.nih.gov

Insert Length: 795 Std Error: 0.00

Seq primer: -40m13 fwd. ET from Amerham

High quality sequence scop: 301.

Location/Qualifiers

FEATURES

source

1..442
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:1427017"
 /tissue_type="substantia nigra"
 /lab_host="DH10B"
 /clone.lib="NCI_CGAP CNS1"
 /note="Organ: Brain; Vector: pCMV-SPORT4; Site_1: SalI;
 Site_2: NotI; Cloned unidirectionally. Primer: Oligo dT.
 Average insert size 1.0 kb."

ORIGIN

Alignment Scores:

Pred. No.:	0.0506	Length:	442
Score:	70.00	Matches:	14
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	1	Gaps:	0

US-09-017-715A-2_COPY_94_107 (1-14) x AA804675 (1-442)

Qy 1 ValValArgLySGluAspLeuArgProSerAlaProGlnGln 14
 |||||
 Db 391 GTGGTGGCGCAGAGAGACTTGAGGCGCATCTGCCCCCAACAG 350

RESULT 13

BG826435 451 bp mRNA linear EST 22-MAY-2001
 LOCUS BG826435
 DEFINITION 602750062F1 NTH_MGC_17 Homo sapiens cDNA clone IMAGE:4903046 5',
 mRNA sequence.

ACCESSION BG826435

VERSION BG826435.1 GI:14174022

KEYWORDS EST.

SOURCE Homo sapiens (human)
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 451)
AUTHORS NIH-MGC <http://mgc.ncl.nih.gov/>.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapsb-remail.nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Ling Hong/Rubin Laboratory
DNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at: <http://image.llnl.gov>
Plate: L1CM1800 row: K column: 15
High quality sequence stop: 447.
Location/Qualifiers
1..451
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:4903046"
/lab_host="DH10B (phage-resistant)"
/clone_id="NIH_MGC_17"
/note="Organ: muscle; Vector: pOTB7; Site: 1: EcoRI; Site 2: XhoI; cDNA made by oligo-dT priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGCACGAG(G). Size-selected >500bp for average insert size 1.8kb. Library constructed by Ling Hong in the Laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies)."

ORIGIN
Alignment Scores:
Pred. No.: 0.0517 Length: 451
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservatve: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-09-017-715a-2_COPY_94_107 (1-14) x BG826435 (1-451)
Cy 1 ValValArgLySGUApLeuArgProSerAlaProGInGln 14
Db 349 GTGGTGGCAAGAGGACCTTGAGCCATGCCCCCAACAG 390

RESULT 14
BX474511 462 bp mRNA linear EST 04-SEP-2003
LOCUS DKFZ686824170.1 686 (synonym: hlcc3) Homo sapiens CDNA clone
DEFINITION DKFZ686824170.5', mRNA sequence.
ACCESSION BX474511
VERSION BX474511.1 GI:31668718
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 462)
AUTHORS Bloeker, H., Boecher, M., Mewes, H.W., Weil, B., Amid, C., Osaenger, A., Robo, G., Han, M. and Wiemann, S.
TITLE (Bloeker, H., Boecher, M., Mewes, H.W., Weil, B., Amid, C., et al.)
JOURNAL Unpublished (2003)
COMMENT Contact: MIPS
MIPS Ingolstaedter Landstr.1, D-85764 Neuberg, Germany
This is the 5' sequence of the clone insert

FEATURES
source
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/clone_id="686 (synonym: hlcc3)"
/note="Vector: pTriplex2; Site: 1: SfiIA; Site: 2: SfiIB; CDNA collection"

Clone from S. Wiemann, Molecular Genome Analysis, German Cancer Research Center (DKFZ); Email: s.wiemann@dkfz-heidelberg.de, sequenced by GBR (National Research Centre for Biotechnology Ltd., Braunschweig/Germany) within the CDNA sequencing consortium of the German Genome Project.
No sl sequence available.
This clone (DKFZ686824170) is available at the RZPD in Berlin. Please contact the RZPD: Resourcenzentrum, Heubnerweg 6, 14059 Berlin-Charlottenburg, GERMANY; Email: clone@rzpd.de.

FEATURES
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/clone="DKFZ686824170"
/dev_stage="adult"
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/clone_id="686 (synonym: hlcc3)"
/note="Vector: pTriplex2; Site: 1: SfiIA; Site: 2: SfiIB; CDNA collection"

ORIGIN
Alignment Scores:
Pred. No.: 0.0531 Length: 462
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservatve: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 5 Gaps: 0

US-09-017-715a-2_COPY_94_107 (1-14) x BX474511 (1-462)
Cy 1 ValValArgLySGUApLeuArgProSerAlaProGInGln 14
Db 390 GTGGTGGCAAGAGGACCTTGAGCCATGCCCCCAACAG 431

RESULT 15
CB107161 467 bp mRNA linear EST 28-JAN-2003
LOCUS CB107161
DEFINITION K-EST0145400 L3SN0475 Homo sapiens CDNA clone L3SN0475-14-A11 5', mRNA sequence.
ACCESSION CB107161
VERSION CB107161.1 GI:27932968
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 467)
AUTHORS Kim, N.S., Hahn, Y., Oh, J.H., Lee, J.Y., Ahn, H.Y., Chu, M.Y., Kim, M.R., Oh, K.J., Cheong, J.E., Sohn, H.Y., Kim, D.M., Park, H.S., Kim, S. and Kim, Y.S.
TITLE 21C Frontier Korean EST Project 2001
JOURNAL Unpublished (2002)
COMMENT Contact: Kim YS
Genome Research Center
Korea Research Institute of Bioscience & Biotechnology
52 Boeun-dong Yuseong-gu, Daejeon 305-333, South Korea
Tel: +82-42-860-4470
Fax: +82-42-860-4409
Email: yongseung@mail.kribb.re.kr
Plate: 14 row: A column: 11
High quality sequence stop: 467.
Location/Qualifiers
1..467
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="L3SN0475-14-A11"
/sex="M"
/cell_line="SNU-475"
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/clone_id="L3SN0475"

GenCore version 5.1.6
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OM protein - nucleic search, using frame_plus model

Run on: May 4, 2005, 10:17:21 ; Search time 113.879 Seconds
(without alignments)
748.404 Million cell updates/sec

Title: US-09-017-715a-2_COPY_94_107
Perfect score: 70
Sequence: 1 VAKEDRPSAPQ 14

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Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 5642217 seqs, 3043843248 residues
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Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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Database :
Published Applications NA:
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10: /cgn2_6/ptodata/1/pubpna/US09B_PUBCOMB.seq:*
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12: /cgn2_6/ptodata/1/pubpna/US09_NEW_PUB.seq:*
13: /cgn2_6/ptodata/1/pubpna/US10_PUBCOMB.seq:*
14: /cgn2_6/ptodata/1/pubpna/US10B_PUBCOMB.seq:*
15: /cgn2_6/ptodata/1/pubpna/US10C_PUBCOMB.seq:*
16: /cgn2_6/ptodata/1/pubpna/US10D_PUBCOMB.seq:*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	70	100.0	384	19 US-10-826-157-5	Sequence 5, Appl
2	70	100.0	479	10 US-09-918-995-2705	Sequence 2705, Appl
3	70	100.0	550	9 US-09-954-531-613	Sequence 613, Appl
4	70	100.0	550	17 US-10-453-478-1	Sequence 1, Appl
5	70	100.0	550	17 US-10-843-641-1680	Sequence 1680, Appl
6	70	100.0	720	14 US-10-097-340-297	Sequence 297, Appl
7	70	100.0	720	14 US-10-282-174-469	Sequence 469, Appl
8	70	100.0	720	19 US-10-600-009-469	Sequence 171, Appl
9	70	100.0	796	9 US-09-925-298-171	Sequence 171, Appl
10	70	100.0	796	14 US-10-102-806-171	Sequence 172, Appl
11	53	75.7	478	9 US-09-925-298-172	Sequence 172, Appl
12	53	75.7	478	14 US-10-102-806-172	Sequence 388, Appl
13	53	75.7	4606	17 US-10-440-425-388	Sequence 72, Appl
14	53	75.7	5666	17 US-10-282-174-72	Sequence 72, Appl
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16	53	75.7	5666	19 US-10-600-009-72	Sequence 73, Appl
17	53	75.7	5666	19 US-10-600-009-73	Sequence 483, Appl
18	53	75.7	6012	19 US-10-282-174-483	Sequence 483, Appl
19	53	75.7	6012	19 US-10-600-009-483	Sequence 1, Appl
20	48	68.6	135638	16 US-10-314-657-1	Sequence 1, Appl
21	48	68.6	135638	19 US-10-473-193-1	Sequence 23678, A
22	47	67.1	657	17 US-10-282-122A-22679	Sequence 165876, A
23	46	65.7	794	18 US-10-425-115-165876	Sequence 141921, A
24	45	64.3	435	18 US-10-425-115-141921	Sequence 6012, Ap
25	45	64.3	465	9 US-09-864-761-6012	Sequence 101237, A
26	45	64.3	1077	13 US-10-027-632-101236	Sequence 101237, A
27	45	64.3	1077	13 US-10-027-632-101237	Sequence 101237, A
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30	45	64.3	1435	17 US-10-282-122A-26255	Sequence 26255, A
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32	45	64.3	1473	17 US-10-282-122A-28635	Sequence 28635, A
33	45	64.3	1488	9 US-09-738-626-1425	Sequence 1425, Ap
34	45	64.3	1491	17 US-10-282-122A-17537	Sequence 17537, A
35	45	64.3	16010	17 US-10-085-117-262	Sequence 262, Appl
36	45	64.3	3309400	9 US-09-738-626-1	Sequence 1, Appl
37	44	62.9	395	18 US-10-425-115-118311	Sequence 118311, A
38	44	62.9	661	15 US-10-259-165-660	Sequence 660, Appl
39	44	62.9	775	17 US-10-424-599-9676	Sequence 9676, Ap
40	44	61.4	412	18 US-10-425-115-52591	Sequence 52591, A
41	43	61.4	465	9 US-09-864-761-4889	Sequence 4889, Ap
42	43	61.4	547	16 US-10-029-386-9553	Sequence 9553, Ap
43	43	61.4	571	9 US-09-864-761-9722	Sequence 9722, Ap
44	43	61.4	800	18 US-10-363-345A-37659	Sequence 37659, A
45	43	61.4	800	18 US-10-363-345A-37660	Sequence 37660, A

ALIGNMENTS

RESULT 1
US-10-826-157-5
Sequence 5, Application US/10826157
Publication No. US20050064548A1
GENERAL INFORMATION:
APPLICANT: Lindquist, Susan L.
TITLE OF INVENTION: YEAST ECTOPICALLY EXPRESSING ABNORMALLY PROCESSED PROTEINS AND USES THEREFOR
FILE REFERENCE: 17481-003001
CURRENT APPLICATION NUMBER: US/10/826,157
CURRENT FILING DATE: 2004-04-16
PRIOR APPLICATION NUMBER: US 60/472,317
PRIOR FILING DATE: 2003-05-20
PRIOR APPLICATION NUMBER: US 60/463,284
PRIOR FILING DATE: 2003-04-16
NUMBER OF SEQ ID NOS: 8
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 5
LENGTH: 384

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/ TYPE: DNA
/ ORGANISM: Homo sapiens
US-10-826-157-5
Alignment Scores:
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Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 19 Gaps: 0

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QY 1 ValValArgLysGluAspLeuArgProSerAlaProGlnGln 14
Db 280 GTGGTGGCGAAGAGGACTTGAGGCCATCTGCCCCCAACAG 321

RESULT 2
US-09-918-995-2705
/ Sequence 2705, Application US/09918995
/ Publication No. US20030073623A1
/ GENERAL INFORMATION:
/ APPLICANT: HySeq, Inc.
/ TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
/ FILE OF INVENTION: FROM VARIOUS CDNA LIBRARIES
/ FILE REFERENCE: 20411-756
/ CURRENT APPLICATION NUMBER: US/09/918,995
/ CURRENT FILING DATE: 2001-07-30
/ PRIOR APPLICATION NUMBER: US/09/235,076
/ PRIOR FILING DATE: 1999-01-20
/ NUMBER OF SEQ ID NOS: 38054
/ SOFTWARE: FastSeq for Windows Version 3.0
/ SEQ ID NO 2705
/ LENGTH: 479
/ TYPE: DNA
/ ORGANISM: Homo sapiens
/ FEATURE:
/ NAME/KEY: misc.feature
/ LOCATION: (1)...(479)
/ OTHER INFORMATION: n = A,T,C or G
US-09-918-995-2705

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Pred. No.: 0.00193 Length: 479
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 10 Gaps: 0

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QY 1 ValValArgLysGluAspLeuArgProSerAlaProGlnGln 14
Db 293 GTGGTGGCGAAGAGGACTTGAGGCCATCTGCCCCCAACAG 334

RESULT 3
US-09-954-531-613
/ Sequence 613, Application US/09954531
/ Patent No. US20020165180A1
/ GENERAL INFORMATION:
/ APPLICANT: Weaver, Zoe
/ TITLE OF INVENTION: Process for Identifying Anti-Cancer Therapeutic Agents Using Canc
/ FILE REFERENCE: 689290-77
/ CURRENT APPLICATION NUMBER: US/09/954,531
/ CURRENT FILING DATE: 2002-05-02
/ PRIOR APPLICATION NUMBER: US/60/233,133
/ PRIOR FILING DATE: 2000-09-18
/ PRIOR APPLICATION NUMBER: US/60/234,009
/ PRIOR FILING DATE: 2000-09-20
/ PRIOR APPLICATION NUMBER: US/60/234,034
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/ PRIOR FILING DATE: 2000-09-20
/ PRIOR APPLICATION NUMBER: US/60/234,509
/ PRIOR FILING DATE: 2000-09-22
/ PRIOR APPLICATION NUMBER: US/60/234,567
/ PRIOR FILING DATE: 2000-09-22
/ NUMBER OF SEQ ID NOS: 1392
/ SOFTWARE: PatentIn version 3.0
/ SEQ ID NO 613
/ LENGTH: 550
/ TYPE: DNA
/ ORGANISM: Homo sapiens
US-09-954-531-613

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Pred. No.: 0.00222 Length: 550
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x US-09-954-531-613 (1-550)
QY 1 ValValArgLysGluAspLeuArgProSerAlaProGlnGln 14
Db 291 GTGGTGGCGAAGAGGACTTGAGGCCATCTGCCCCCAACAG 332

RESULT 4
US-10-453-478-1
/ Sequence 1, Application US/10453478
/ Publication No. US20030208043A1
/ GENERAL INFORMATION:
/ APPLICANT: Paul Moore, Reiner Gentz, Hongjin Ji,
/ TITLE OF INVENTION: Human Genes, Sequences and
/ EXPRESSION PRODUCTS
/ NUMBER OF SEQUENCES: 22
/ CORRESPONDENCE ADDRESS:
/ ADDRESSEE: CARELLA, BYRNE, BAIN, GILFILLAN,
/ STREET: 6 BECKER FARM ROAD
/ CITY: ROSELAND
/ STATE: NEW JERSEY
/ COUNTRY: USA
/ ZIP: 07068
/ COMPUTER READABLE FORM:
/ MEDIUM TYPE: 3.5 INCH DISKETTE
/ COMPUTER: IBM PS/2
/ OPERATING SYSTEM: MS-DOS
/ SOFTWARE: WORD PERFECT 5.1
/ CURRENT APPLICATION DATA:
/ APPLICATION NUMBER: US/10/453,478
/ FILING DATE: 04-Jun-2003
/ CLASSIFICATION: 536
/ PRIOR APPLICATION DATA:
/ APPLICATION NUMBER: US/08/705,771
/ FILING DATE: August 30, 1996
/ ATTORNEY/AGENT INFORMATION:
/ NAME: MULLINS, J.G.
/ REGISTRATION NUMBER: 33,073
/ REFERENCE/DOCKET NUMBER: 325800-346 (PFI96)
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: 973-994-1700
/ TELEFAX: 973-994-1744
/ INFORMATION FOR SEQ ID NO: 1:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 550 base pairs
/ TYPE: nucleic acid
/ STRANDEDNESS: single
/ TOPOLOGY: linear
/ MOLECULE TYPE: DNA
/ SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-10-453-478-1
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Alignment Scores:
Pred. No.: 0.00222 Length: 550
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 17 Gaps: 0

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Db 291 GTGCTGCCAAGAGAGACTTGAGGCCATCTCCCCCAACAG 332

RESULT 5
US-10-843-641A-1680
; Sequence 1680, Application US/10843641A
; Publication No. US20050064454A1
; GENERAL INFORMATION:
; APPLICANT: Avalon Pharmaceuticals, Inc.
; TITLE OF INVENTION: Cancer Gene Determination and Therapeutic Screening Using
; FILE REFERENCE: 689290-189
; CURRENT APPLICATION NUMBER: US/10/843,641A
; CURRENT FILING DATE: 2004-05-12
; PRIOR APPLICATION NUMBER: US/09/873,367
; PRIOR FILING DATE: 2001-06-05
; PRIOR APPLICATION NUMBER: US/09/954,531
; PRIOR FILING DATE: 2001-09-18
; PRIOR APPLICATION NUMBER: US/09/954,456
; PRIOR FILING DATE: 2001-09-25
; PRIOR APPLICATION NUMBER: US/09/962,436
; PRIOR FILING DATE: 2001-09-25
; PRIOR APPLICATION NUMBER: US/09/962,832
; PRIOR FILING DATE: 2001-09-25
; PRIOR APPLICATION NUMBER: US/09/964,824
; PRIOR FILING DATE: 2001-09-27
; PRIOR APPLICATION NUMBER: US/09/967,768
; PRIOR FILING DATE: 2001-09-28
; PRIOR APPLICATION NUMBER: US/09/968,007
; PRIOR FILING DATE: 2001-10-02
; PRIOR APPLICATION NUMBER: US/09/969,347
; PRIOR FILING DATE: 2001-10-02
; PRIOR APPLICATION NUMBER: US/09/969,708
; PRIOR FILING DATE: 2001-10-03
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 8447
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 1680
; LENGTH: 550
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-843-641A-1680

Alignment Scores:
Pred. No.: 0.00222 Length: 550
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 19 Gaps: 0

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Db 291 GTGCTGCCAAGAGAGACTTGAGGCCATCTCCCCCAACAG 332

RESULT 6
US-10-097-340-297
; Sequence 297, Application US/10097340
; Publication No. US20030087250A1

GENERAL INFORMATION:
; APPLICANT: John MONAHAN
; APPLICANT: Manjula GANNAVARAPU
; APPLICANT: Sebastian HOERSCHE
; APPLICANT: Shubhangi KAMATKAR
; APPLICANT: Steve G. KOVATS
; APPLICANT: Rachel E. MEYERS
; APPLICANT: Michael MORRISSEY
; APPLICANT: Peter OLANDT
; APPLICANT: Ami SEN
; APPLICANT: Peter VEIBY
; APPLICANT: Gordon B. MILLS
; APPLICANT: Robert C. BAST, Jr.
; APPLICANT: Karen LU
; APPLICANT: Rosemarie SCHMANDT
; APPLICANT: Xumei ZHAO
; APPLICANT: Karen GLATT
; TITLE OF INVENTION: Nucleic Acid Molecules and Proteins For The Identification,
; TITLE OF INVENTION: Assessment, Prevention, and Therapy of Ovarian Cancer
; FILE REFERENCE: MRI-030
; CURRENT APPLICATION NUMBER: US/10/097,340
; CURRENT FILING DATE: 2002-03-14
; PRIOR APPLICATION NUMBER: 60/276,025
; PRIOR FILING DATE: 2001-03-14
; PRIOR APPLICATION NUMBER: 60/325,149
; PRIOR FILING DATE: 2001-09-26
; PRIOR APPLICATION NUMBER: 60/276,026
; PRIOR FILING DATE: 2001-03-14
; PRIOR APPLICATION NUMBER: 60/324,967
; PRIOR FILING DATE: 2001/09/26
; PRIOR APPLICATION NUMBER: 60/311,732
; PRIOR FILING DATE: 2001-08-10
; PRIOR APPLICATION NUMBER: 60/325,102
; PRIOR FILING DATE: 2001-09-26
; PRIOR APPLICATION NUMBER: 60/323,580
; PRIOR FILING DATE: 2001-09-19
; NUMBER OF SEQ ID NOS: 363
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 297
; LENGTH: 720
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-097-340-297

Alignment Scores:
Pred. No.: 0.00289 Length: 720
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 14 Gaps: 0

US-09-017-715a-2_COPY_94_107 (1-14) x US-10-097-340-297 (1-720)
Qy 1 ValValAArgLySGUAAPLeuArgProSeRaLaProGInGIn 14
Db 328 GTGCTGCCAAGAGAGACTTGAGGCCATCTCCCCCAACAG 369

RESULT 7
US-10-282-174-469
; Sequence 469, Application US/10282174
; Publication No. US20030224380A1
; GENERAL INFORMATION:
; APPLICANT: Becker, Kenneth David
; APPLICANT: Velicelebi, Gonul
; APPLICANT: Elliot, Kathryn J.
; APPLICANT: Wang, Xin
; APPLICANT: Tanzi, Rudolph E.
; APPLICANT: Bertam, Lars
; APPLICANT: Saunders, Aleister J.
; APPLICANT: Mullin, Kristina M.
; APPLICANT: Sampson, Andrew Johnson
; APPLICANT: Blacker, Deborah Lynne
```

TITLE OF INVENTION: GENES AND POLYMORPHISMS ON CHROMOSOME 10
TITLE OF INVENTION: ASSOCIATED WITH ALZHEIMER'S DISEASE AND OTHER
FILE REFERENCE: 37481-3308
CURRENT APPLICATION NUMBER: US 10/282,174
CURRENT FILING DATE: 2002-10-25
PRIOR APPLICATION NUMBER: US 60/339,525
PRIOR FILING DATE: 2001-10-25
PRIOR APPLICATION NUMBER: US 60/338,010
PRIOR FILING DATE: 2001-11-08
PRIOR APPLICATION NUMBER: US 60/336,929
PRIOR FILING DATE: 2001-11-08
PRIOR APPLICATION NUMBER: US 60/338,363
PRIOR FILING DATE: 2001-11-09
PRIOR APPLICATION NUMBER: US 60/337,052
PRIOR FILING DATE: 2001-12-04
PRIOR APPLICATION NUMBER: US 60/368,919
PRIOR FILING DATE: 2002-03-28
NUMBER OF SEQ ID NOS: 564
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 469
LENGTH: 720
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: allele
LOCATION: 30,57,85,243,250,377,512,531,555,561,672
OTHER INFORMATION: N is any
US-10-282-174-469
Alignment Scores:
Pred. No.: 0.00289 Length: 720
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
Gaps: 0
DB: 17
US-09-017-715a-2_copy_94_107 (1-14) x US-10-282-174-469 (1-720)
Qy 1 ValValAArgLyGluAspLeuArgProSerAlaProGlnGln 14
Db 328 GTGGTGGCGCAGAGAGACTTGAGGCCATCTGCCCCCAACAG 369
RESULT 8
US-10-600-009-469
Sequence 469, Application US/10600009
Publication No. US2005009031A1
GENERAL INFORMATION:
APPLICANT: Becker, Kenneth David
APPLICANT: Velicelbi, Gonul
APPLICANT: Elliot, Kathryn J.
APPLICANT: Wang, Xin
APPLICANT: Tanzi, Rudolph E.
APPLICANT: Bertram, Lars
APPLICANT: Saunders, Aleister J.
APPLICANT: Mullin, Kristina M.
APPLICANT: Sampson, Andrew Johnson
APPLICANT: Blacker, Deborah Lynne
TITLE OF INVENTION: GENES AND POLYMORPHISMS ON CHROMOSOME 10
TITLE OF INVENTION: ASSOCIATED WITH ALZHEIMER'S DISEASE AND OTHER
FILE REFERENCE: 37481-3308
CURRENT APPLICATION NUMBER: US/10/600,009
CURRENT FILING DATE: 2003-06-18
PRIOR APPLICATION NUMBER: US 60/339,525
PRIOR FILING DATE: 2001-10-25
PRIOR APPLICATION NUMBER: US 60/338,010
PRIOR FILING DATE: 2001-11-08
PRIOR APPLICATION NUMBER: US 60/336,929
PRIOR FILING DATE: 2001-11-08
PRIOR APPLICATION NUMBER: US 60/338,363
PRIOR FILING DATE: 2001-11-09

PRIOR APPLICATION NUMBER: US 60/337,052
PRIOR FILING DATE: 2001-12-04
PRIOR APPLICATION NUMBER: US 60/368,919
PRIOR FILING DATE: 2002-03-28
PRIOR APPLICATION NUMBER: US 10/282,174
PRIOR FILING DATE: 2002-10-25
NUMBER OF SEQ ID NOS: 564
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 469
LENGTH: 720
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: allele
LOCATION: 30,57,85,243,250,377,512,531,555,561,672
OTHER INFORMATION: N is any
US-10-600-009-469
Alignment Scores:
Pred. No.: 0.00289 Length: 720
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
Gaps: 0
DB: 19
US-09-017-715a-2_copy_94_107 (1-14) x US-10-600-009-469 (1-720)
Qy 1 ValValAArgLyGluAspLeuArgProSerAlaProGlnGln 14
Db 328 GTGGTGGCGCAGAGAGACTTGAGGCCATCTGCCCCCAACAG 369
RESULT 9
US-09-925-298-171
Sequence 171, Application US/09925298
Publication No. US20020039764A1
GENERAL INFORMATION:
APPLICANT: Rosen et al.
TITLE OF INVENTION: Nucleic Acids, Proteins and Antibodies
FILE REFERENCE: PA103
CURRENT APPLICATION NUMBER: US/09/925,298
CURRENT FILING DATE: 2001-08-10
PRIOR APPLICATION NUMBER: PCT/US00/05881
PRIOR FILING DATE: 2000-03-08
PRIOR APPLICATION NUMBER: 60/124,270
PRIOR FILING DATE: 1998-03-12
NUMBER OF SEQ ID NOS: 846
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 171
LENGTH: 796
TYPE: DNA
ORGANISM: Homo sapiens
US-09-925-298-171
Alignment Scores:
Pred. No.: 0.00319 Length: 796
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
Gaps: 0
DB: 9
US-09-017-715a-2_copy_94_107 (1-14) x US-09-925-298-171 (1-796)
Qy 1 ValValAArgLyGluAspLeuArgProSerAlaProGlnGln 14
Db 388 GTGGTGGCGCAGAGAGACTTGAGGCCATCTGCCCCCAACAG 429
RESULT 10
US-10-102-806-171
Sequence 171, Application US/10102806
Publication No. US20030054421A1
GENERAL INFORMATION:


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; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins and Antibodies
; FILE REFERENCE: PA103P1C1
; CURRENT APPLICATION NUMBER: US/10/102,806
; PRIOR FILING DATE: 2002-03-22
; PRIOR APPLICATION NUMBER: 09/925,298
; PRIOR FILING DATE: 2001-08-10
; PRIOR APPLICATION NUMBER: PCT/US00/05881
; PRIOR FILING DATE: 2000-03-08
; PRIOR APPLICATION NUMBER: 60/124,270
; PRIOR FILING DATE: 1999-03-12
; NUMBER OF SEQ ID NOS: 846
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 171
; LENGTH: 796
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-10-102-806-171

Alignment Scores:
Pred. No.: 0.00319      Length: 796
Score: 70.00           Matches: 14
Percent Similarity: 100.00%  Conservative: 0
Best Local Similarity: 100.00%  Mismatches: 0
Query Match: 14        Indels: 0
DB: 14                Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x US-10-102-806-171 (1-796)

Cy 1 ValValAlrGlyGluApLeuArgProSerAlaProGlnGln 14
Db 388 GTGCTGCCGACAGAGGACTTGGCCCATCTCCCCCAACAG 429

RESULT 11
US-09-925-298-172
; Sequence 172, Application US/09925298
; Publication No. US20020039764A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins and Antibodies
; FILE REFERENCE: PA103
; CURRENT APPLICATION NUMBER: US/09/925,298
; CURRENT FILING DATE: 2001-08-10
; PRIOR APPLICATION NUMBER: PCT/US00/05881
; PRIOR FILING DATE: 2000-03-08
; PRIOR APPLICATION NUMBER: 60/124,270
; PRIOR FILING DATE: 1999-03-12
; NUMBER OF SEQ ID NOS: 846
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 172
; LENGTH: 478
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-925-298-172

Alignment Scores:
Pred. No.: 2.27      Length: 478
Score: 53.00         Matches: 10
Percent Similarity: 100.00%  Conservative: 1
Best Local Similarity: 90.91%  Mismatches: 0
Query Match: 75.71%  Indels: 0
DB: 9                Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x US-09-925-298-172 (1-478)

Cy 4 LyeGluApLeuArgProSerAlaProGlnGln 14
Db 80 CAGAGAGACTTGGAGCCATCTGCCCAACAG 112

RESULT 12
US-10-102-806-172
; Sequence 172, Application US/10102806
; Publication No. US20030054421A1
```

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; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins and Antibodies
; FILE REFERENCE: PA103P1C1
; CURRENT APPLICATION NUMBER: US/10/102,806
; PRIOR FILING DATE: 2002-03-22
; PRIOR APPLICATION NUMBER: 09/925,298
; PRIOR FILING DATE: 2001-08-10
; PRIOR APPLICATION NUMBER: PCT/US00/05881
; PRIOR FILING DATE: 2000-03-08
; PRIOR APPLICATION NUMBER: 60/124,270
; PRIOR FILING DATE: 1999-03-12
; NUMBER OF SEQ ID NOS: 846
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 172
; LENGTH: 478
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-10-102-806-172

Alignment Scores:
Pred. No.: 2.27      Length: 478
Score: 53.00         Matches: 10
Percent Similarity: 100.00%  Conservative: 1
Best Local Similarity: 90.91%  Mismatches: 0
Query Match: 75.71%  Indels: 0
DB: 14                Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x US-10-102-806-172 (1-478)

Cy 4 LyeGluApLeuArgProSerAlaProGlnGln 14
Db 80 CAGAGAGACTTGGAGCCATCTGCCCAACAG 112

RESULT 13
US-10-240-425-388
; Sequence 388, Application US/10240425
; Publication No. US2004003502A1
; GENERAL INFORMATION:
; APPLICANT: Williams, Amanda
; APPLICANT: Boland, Joseph F.
; APPLICANT: Lord, Reginald V.
; APPLICANT: Alvarez, Chris
; APPLICANT: Wetzel, Jon C.
; APPLICANT: Schett, Uwe
; APPLICANT: Vockley, Joseph G.
; TITLE OF INVENTION: Gene Expression Profiles in Esophageal Tissue
; FILE REFERENCE: 44921-5026
; CURRENT APPLICATION NUMBER: US/10/240,425
; CURRENT FILING DATE: 2002-09-30
; PRIOR APPLICATION NUMBER: PCT/US01/09847
; PRIOR FILING DATE: 2001-03-28
; PRIOR APPLICATION NUMBER: US 60/193,446
; PRIOR FILING DATE: 2000-03-31
; NUMBER OF SEQ ID NOS: 1588
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 388
; LENGTH: 4606
; TYPE: DNA
; ORGANISM: Homo sapiens
; OTHER INFORMATION: Genbank Accession No. US2004003502A1 AF044311
; US-10-240-425-388

Alignment Scores:
Pred. No.: 21.1      Length: 4606
Score: 53.00         Matches: 10
Percent Similarity: 100.00%  Conservative: 1
Best Local Similarity: 90.91%  Mismatches: 0
Query Match: 75.71%  Indels: 0
DB: 17                Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x US-10-240-425-388 (1-4606)
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Qy 4 LysGluAspLeuArgProSerAlaProGlnGln 14
Db 3950 CAGGAGACTTGAAGCCATCTGCCCCCAACAG 3982

RESULT 14
US-10-282-174-72
Sequence 72, Application US/10282174
Publication No. US20030224380A1
GENERAL INFORMATION:
APPLICANT: Becker, Kenneth David
APPLICANT: Velicelebi, Goni
APPLICANT: Elliot, Kathryn J.
APPLICANT: Wang, Xin
APPLICANT: Tanzi, Rudolph E.
APPLICANT: Bertam, Lars
APPLICANT: Saunders, Aleister J.
APPLICANT: Sampson, Andrew Johnson
APPLICANT: Blacker, Deborah Lynne
TITLE OF INVENTION: GENES AND POLYMORPHISMS ON CHROMOSOME 10
TITLE OF INVENTION: ASSOCIATED WITH ALZHEIMER'S DISEASE AND OTHER
TITLE OF INVENTION: NEURODEGENERATIVE DISEASES
FILE REFERENCE: 37481-3308
CURRENT FILING DATE: 2002-10-25
PRIOR FILING DATE: 2001-10-25
PRIOR FILING DATE: 2001-10-25
PRIOR FILING DATE: 2001-10-25
PRIOR FILING DATE: 2001-11-08
PRIOR FILING DATE: 2001-11-08
PRIOR FILING DATE: 2001-11-08
PRIOR FILING DATE: 2001-11-08
PRIOR FILING DATE: 2001-11-08
PRIOR FILING DATE: 2001-11-09
PRIOR FILING DATE: 2001-12-04
PRIOR FILING DATE: 2001-12-04
PRIOR FILING DATE: 2002-03-28
NUMBER OF SEQ ID NOS: 564
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 72
LENGTH: 5666
TYPE: DNA
ORGANISM: Homo sapiens
US-10-282-174-72

Alignment Scores:
Pred. No.: 25.9
Score: 53.00
Percent Similarity: 100.00%
Best Local Similarity: 90.91%
Query Match: 75.71%
DB: 17
Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x US-10-282-174-72 (1-5666)
Qy 4 LysGluAspLeuArgProSerAlaProGlnGln 14
Db 4512 CAGGAGACTTGAAGCCATCTGCCCCCAACAG 4544

RESULT 15
US-10-282-174-73
Sequence 73, Application US/10282174
Publication No. US20030224380A1
GENERAL INFORMATION:
APPLICANT: Becker, Kenneth David
APPLICANT: Velicelebi, Goni
APPLICANT: Elliot, Kathryn J.
APPLICANT: Wang, Xin
APPLICANT: Tanzi, Rudolph E.
APPLICANT: Bertam, Lars
APPLICANT: Saunders, Aleister J.
APPLICANT: Mullin, Kristina M.
```

```
APPLICANT: Sampson, Andrew Johnson
APPLICANT: Blacker, Deborah Lynne
TITLE OF INVENTION: GENES AND POLYMORPHISMS ON CHROMOSOME 10
TITLE OF INVENTION: ASSOCIATED WITH ALZHEIMER'S DISEASE AND OTHER
TITLE OF INVENTION: NEURODEGENERATIVE DISEASES
FILE REFERENCE: 37481-3308
CURRENT FILING DATE: 2002-10-25
PRIOR FILING DATE: 2002-10-25
PRIOR FILING DATE: 2001-10-25
PRIOR FILING DATE: 2001-10-25
PRIOR FILING DATE: 2001-11-08
PRIOR FILING DATE: 2001-11-08
PRIOR FILING DATE: 2001-11-08
PRIOR FILING DATE: 2001-11-08
PRIOR FILING DATE: 2001-11-09
PRIOR FILING DATE: 2001-12-04
PRIOR FILING DATE: 2001-12-04
PRIOR FILING DATE: 2002-03-28
NUMBER OF SEQ ID NOS: 564
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 73
LENGTH: 5666
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: allele
LOCATION: 560,590,617,645,915,987,1723,1943,1950,3151,3178,3189,3284,
LOCATION: 4276,4311,4552,4995,5019,5025,5112,5136,5421,5648,5517
OTHER INFORMATION: N is any
FEATURE:
NAME/KEY: allele
LOCATION: 3779
OTHER INFORMATION: deletion: T
FEATURE:
NAME/KEY: allele
LOCATION: 4156
OTHER INFORMATION: insertion following nucleotide 4155
NAME/KEY: allele
LOCATION: 4976
OTHER INFORMATION: deletion: C
US-10-282-174-73

Alignment Scores:
Pred. No.: 25.9
Score: 53.00
Percent Similarity: 100.00%
Best Local Similarity: 90.91%
Query Match: 75.71%
DB: 17
Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x US-10-282-174-73 (1-5666)
Qy 4 LysGluAspLeuArgProSerAlaProGlnGln 14
Db 4512 CAGGAGACTTGAAGCCATCTGCCCCCAACAG 4544

Search completed: May 4, 2005, 16:39:43
Job time : 126.546 secs
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GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM protein - nucleic search, using frame_plus.p2n model

Run on: May 4, 2005, 09:07:55 ; Search time 32.4161 Seconds

(without alignments)

706.682 Million cell updates/sec

Title: US-09-017-715A-2_COPY_94_107

Perfect score: 70

Sequence: 1 VAKKEDRPSAPQ 14

Scoring table:

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Ygapop 10.0 , Ygapext 0.5	
Fgapop 6.0 , Fgapext 7.0	
Delop 6.0 , Delext 7.0	

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Command line parameters:

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-LOOPEXT=0 -UNITS=bits -START=1 -END=-1 -MATRIX=blosum62 -TRANS=human40.cdi
-LIST=45 -DOCALLIGN=200 -THR SCORE=pct -THR MAX=100 -THR MIN=0 -ALIGN=15
-MODE=LOCAL -OUTFMT=pct -NOM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000
-USRR=US09017715 -OCGN 1 1 116 -runat_04052005_100745_25632 -NCPU=6 -ICPU=3
-NO MMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG
-DEV TIMEOUT=120 -MAIN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

Issued_Patents_NA.*
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2: /cgn2_6/ptodata/1/ina/5B.COMB.seq.*
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4: /cgn2_6/ptodata/1/ina/6B.COMB.seq.*
5: /cgn2_6/ptodata/1/ina/PTCUS.COMB.seq.*
6: /cgn2_6/ptodata/1/ina/backfill1.seq.*

Print. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed.
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	70	100.0	550	3	US-08-705-771-1
2	70	100.0	550	4	US-09-417-540-1
3	70	100.0	702	4	US-09-949-016-1915
4	70	100.0	720	4	US-09-949-016-442
5	53	75.7	8607	4	US-09-949-016-13657
6	53	75.7	8608	4	US-09-949-016-12184
7	47	67.1	825	4	US-09-489-039A-172
8	45	64.3	4403765	3	US-09-103-840A-2
9	45	64.3	4411529	3	US-09-103-840A-1
10	43	61.4	1011	4	US-09-902-540-8804
11	43	61.4	1187	1	US-08-440-856A-2
12	43	61.4	10096	4	US-09-902-540-935

13	42	60.0	689	4	US-09-949-016-5443	Sequence 5443, Ap
14	42	60.0	690	3	US-09-419-568F-24	Sequence 24, Appl
15	42	60.0	690	3	US-09-354-243B-24	Sequence 24, Appl
16	42	60.0	1152	4	US-09-870-574-1	Sequence 1, Appl1
17	42	60.0	4797	3	US-09-419-568F-25	Sequence 25, Appl
18	42	60.0	4797	3	US-09-354-243B-25	Sequence 25, Appl
19	42	60.0	8888	4	US-09-949-016-17185	Sequence 17185, A
20	42	60.0	50950	4	US-09-949-016-16569	Sequence 16569, A
21	41.5	59.3	9005	4	US-09-902-540-7894	Sequence 7894, Ap
22	41.5	59.3	10793	4	US-09-902-540-1062	Sequence 1062, Ap
23	41	58.6	601	4	US-09-949-016-115274	Sequence 115274,
24	41	58.6	601	4	US-09-949-016-144544	Sequence 144544,
25	41	58.6	918	4	US-09-902-540-9560	Sequence 9560, Ap
26	41	58.6	1073	1	US-08-356-405-8	Sequence 8, Appl
27	41	58.6	1074	4	US-08-356-405-8	Sequence 8, Appl
28	41	58.6	1545	4	US-09-270-767-1178	Sequence 1178, Ap
29	41	58.6	1545	4	US-09-270-767-15460	Sequence 15460, A
30	41	58.6	2128	3	US-09-280-116-11	Sequence 11, Appl
31	41	58.6	2217	1	US-07-865-662F-9	Sequence 9, Appl
32	41	58.6	2217	1	US-08-374-219B-9	Sequence 9, Appl
33	41	58.6	2459	3	US-09-443-795-2	Sequence 2, Appl1
34	41	58.6	7386	4	US-09-949-016-13287	Sequence 13287, A
35	41	58.6	8268	1	US-08-375-709-10	Sequence 10, Appl
36	41	58.6	8268	1	US-08-752-929-10	Sequence 10, Appl
37	41	58.6	8268	1	US-09-231-899-83	Sequence 83, Appl
38	41	58.6	15172	4	US-09-902-540-1086	Sequence 1086, Ap
39	41	58.6	15661	4	US-09-949-016-13161	Sequence 13161, A
40	41	58.6	18200	4	US-09-949-016-15660	Sequence 15660, A
41	41	58.6	18200	4	US-09-949-016-15661	Sequence 15661, A
42	41	58.6	19719	4	US-09-949-016-15662	Sequence 15662, A
43	41	58.6	19719	4	US-09-949-016-15663	Sequence 15663, A
44	41	58.6	37895	4	US-08-375-709-1	Sequence 1, Appl1
45	41	58.6	37895	1	US-08-752-929-1	Sequence 1, Appl1

ALIGNMENTS

RESULT 1
US-08-705-771-1
; Sequence 1, Application US/08705771
; Patent No. 6054289
GENERAL INFORMATION:
APPLICANT: Paul Moore, Reiner Genz, Hongjin Ji,
Jian Ni and Jing-Shan Hu
TITLE OF INVENTION: Human Genes, Sequences and
TITLE OF INVENTION: Expression Products
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESSES:
ADDRESSEE: CARELLA, BYRNE, BAIN, GIUFFILIAN,
ADDRESS: CECCHI, STEWART & OLSTEIN
STREET: 6 BECKER FARM ROAD
CITY: ROSELAND
STATE: NEW JERSEY
COUNTRY: USA
ZIP: 07068
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5 INCH DISKETTE
COMPUTER: IBM PS/2
OPERATING SYSTEM: MS-DOS
SOFTWARE: WORD PERFECT 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/705,771
FILING DATE: August 30, 1996
CLASSIFICATION: 536
ATTORNEY/AGENT INFORMATION:
NAME: MULLINS, J.G.
REGISTRATION NUMBER: 33,073
REFERENCE/DOCKET NUMBER: 325800-346 (PT196)
TELECOMMUNICATION INFORMATION:
TELEPHONE: 973-994-1744
TELEFAX: 973-994-1744
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:

LENGTH: 550 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-705-771-1

Alignment Scores:

Pred. No.:	0.00135	Length:	550
Score:	70.00	Matches:	14
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	3	Gaps:	0

US-09-017-715A-2_COPY_94_107 (1-14) x US-08-705-771-1 (1-550)

Qy 1 ValValaArgLyGluAspLeuArgProSerAlaProGlnGln 14
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Db 291 GTGTCGCGAAGAGAGACTTGAGCCATCTGCCCCCAACAG 332

RESULT 2

US-09-417-540-1
Sequence 1, Application US/09417540
Patent No. 6639052

GENERAL INFORMATION:

APPLICANT: Paul Moore, Reiner Gentz, Hongjin Ji,

TITLE OF INVENTION: Human Genes, Sequences and

NUMBER OF SEQUENCES: 22 Expression Products

CORRESPONDENCE ADDRESS:

ADDRESSEE: CARELLA, BYRNE, BAIN, GILFILLAN,

STREET: 6 BECKER FARM ROAD

CITY: ROSELAND

STATE: NEW JERSEY

COUNTRY: USA

ZIP: 07068

MEDIUM TYPE: 3.5 INCH DISKETTE

COMPUTER: IBM PS/2

OPERATING SYSTEM: MS-DOS

SOFTWARE: WORD PERFECT 5.1

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/417,540

FILING DATE: 14-Oct-1999

CLASSIFICATION: <Unknown>

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/705,771

FILING DATE: August 30, 1996

ATTORNEY/AGENT INFORMATION:

NAME: MULLINS, J.G.

REGISTRATION NUMBER: 33,073

REFERENCE/DOCKET NUMBER: 325800-346 (PFI96)

TELECOMMUNICATION INFORMATION:

TELEPHONE: 973-994-1700

TELEFAX: 973-994-1744

INFORMATION FOR SEQ ID NO. 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 550 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA

SEQUENCE DESCRIPTION: SEQ ID NO: 1:

Alignment Scores:

Pred. No.:	0.00135	Length:	550
Score:	70.00	Matches:	14
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0

Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x US-09-417-540-1 (1-550)

Qy 1 ValValaArgLyGluAspLeuArgProSerAlaProGlnGln 14
|||||
Db 291 GTGTCGCGAAGAGAGACTTGAGCCATCTGCCCCCAACAG 332

RESULT 3

US-09-949-016-1915
Sequence 1915, Application US/09949016
Patent No. 6812339

GENERAL INFORMATION:

APPLICANT: VENTER, J. Craig et al.

TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED

FILE REFERENCE: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

CURRENT APPLICATION NUMBER: US/09/949,016

PRIOR FILING DATE: 2000-04-14

PRIOR APPLICATION NUMBER: 60/241,755

PRIOR FILING DATE: 2000-10-20

PRIOR APPLICATION NUMBER: 60/237,768

PRIOR FILING DATE: 2000-10-03

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR FILING DATE: 2000-09-08

NUMBER OF SEQ ID NOS: 207012

SOFTWARE: FastSeq for Windows Version 4.0

SEQ ID NO 1915

LENGTH: 702

TYPE: DNA

ORGANISM: Human

US-09-949-016-1915

Alignment Scores:

Pred. No.:	0.00178	Length:	702
Score:	70.00	Matches:	14
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	4	Gaps:	0

US-09-017-715A-2_COPY_94_107 (1-14) x US-09-949-016-1915 (1-702)

Qy 1 ValValaArgLyGluAspLeuArgProSerAlaProGlnGln 14
|||||
Db 327 GTGTCGCGAAGAGAGACTTGAGCCATCTGCCCCCAACAG 368

RESULT 4

US-09-949-016-442
Sequence 442, Application US/09949016
Patent No. 6812339

GENERAL INFORMATION:
APPLICANT: VENTER, J. Craig et al.

TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED

FILE REFERENCE: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

CURRENT APPLICATION NUMBER: US/09/949,016

PRIOR FILING DATE: 2000-04-14

PRIOR APPLICATION NUMBER: 60/241,755

PRIOR FILING DATE: 2000-10-20

PRIOR APPLICATION NUMBER: 60/237,768

PRIOR FILING DATE: 2000-10-03

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR FILING DATE: 2000-09-08

NUMBER OF SEQ ID NOS: 207012

SOFTWARE: FastSeq for Windows Version 4.0

SEQ ID NO 442

LENGTH: 720

TYPE: DNA

ORGANISM: Human

US-09-949-016-442

Alignment Scores:

Pred. No.: 0.00184 Length: 720
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-09-017-715a-2_COPY_94_107 (1-14) x US-09-949-016-442 (1-720)

QY 1 ValValArgLysGluAspLeuArgProSerAlaProGlnGln 14
|||

Db 328 GTGGTGGCAAGAGGAGCACTTGAGCCATCTGCCCCCAACAG 369

RESULT 5

US-09-949-016-13657
Sequence 13657, Application US/09949016
Patent No. 6812339

GENERAL INFORMATION:

APPLICANT: VENTER, J. Craig et al.

TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED

FILE REFERENCE: CL001307

CURRENT FILING DATE: 2000-04-14

PRIOR APPLICATION NUMBER: 60/241,755

PRIOR FILING DATE: 2000-10-20

PRIOR APPLICATION NUMBER: 60/237,768

PRIOR FILING DATE: 2000-10-03

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR FILING DATE: 2000-09-08

NUMBER OF SEQ ID NOS: 207012

SOFTWARE: FASTSEQ for Windows Version 4.0

SEQ ID NO 13657

LENGTH: 8607

TYPE: DNA

ORGANISM: Human

FEATURE:

NAME/KEY: misc_feature

LOCATION: (1)..(8607)

OTHER INFORMATION: n = A,T,C or G

US-09-949-016-13657

Alignment Scores:

Pred. No.: 31.2 Length: 8607
Score: 53.00 Matches: 10
Percent Similarity: 100.00% Conservative: 1
Best Local Similarity: 90.91% Mismatches: 0
Query Match: 75.71% Indels: 0
DB: 4 Gaps: 0

US-09-017-715a-2_COPY_94_107 (1-14) x US-09-949-016-13657 (1-8607)

QY 4 LysGluAspLeuArgProSerAlaProGlnGln 14
:::|||||

Db 5951 CAGGAGCACTTGAGCCATCTGCCCCCAACAG 5983

RESULT 6

US-09-949-016-12184
Sequence 12184, Application US/09949016
Patent No. 6812339

GENERAL INFORMATION:

APPLICANT: VENTER, J. Craig et al.

TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED

FILE REFERENCE: CL001307

CURRENT FILING DATE: 2000-04-14

PRIOR APPLICATION NUMBER: 60/241,755

PRIOR FILING DATE: 2000-10-20

PRIOR APPLICATION NUMBER: 60/237,768

PRIOR FILING DATE: 2000-10-03

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR FILING DATE: 2000-09-08

NUMBER OF SEQ ID NOS: 207012

SOFTWARE: FASTSEQ for Windows Version 4.0

SEQ ID NO 12184

LENGTH: 8608

TYPE: DNA

ORGANISM: Human

FEATURE:

NAME/KEY: misc_feature

LOCATION: (1)..(8608)

OTHER INFORMATION: n = A,T,C or G

US-09-949-016-12184

QY 3 ArgLysGluAspLeuArgProSerAlaProGlnGln 14
|||

Db 234 CGATTGAGCATCTCGCGCCGCTGGCGCACGCCAG 199

RESULT 8

US-09-103-840A-2/c

Sequence 2, Application US/09103840A

Patent No. 6294328

GENERAL INFORMATION:

APPLICANT: FLEISCHMAN, Robert D.

APPLICANT: WHITE, Owen R.

APPLICANT: PRASER, Claire M.

APPLICANT: VENTER, John C.

TITLE OF INVENTION: DNA SEQUENCES FOR STRAIN ANALYSIS IN MYCOBACTERIUM

FILE REFERENCE: 24366-20007.00

Alignment Scores:

Pred. No.: 31.2 Length: 8608
Score: 53.00 Matches: 10
Percent Similarity: 100.00% Conservative: 1
Best Local Similarity: 90.91% Mismatches: 0
Query Match: 75.71% Indels: 0
DB: 4 Gaps: 0

US-09-017-715a-2_COPY_94_107 (1-14) x US-09-949-016-12184 (1-8608)

QY 4 LysGluAspLeuArgProSerAlaProGlnGln 14
:::|||||

Db 5951 CAGGAGCACTTGAGCCATCTGCCCCCAACAG 5983

RESULT 7

US-09-489-039A-172/c
Sequence 172, Application US/09489039A
Patent No. 6610836

GENERAL INFORMATION:

APPLICANT: GARY Breton et. al

TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA

FILE REFERENCE: 2709.2004001

CURRENT FILING DATE: 2000-01-27

PRIOR APPLICATION NUMBER: US 60/117,747

PRIOR FILING DATE: 1999-01-29

NUMBER OF SEQ ID NOS: 14342

SEQ ID NO 172

LENGTH: 825

TYPE: DNA

ORGANISM: Klebsiella pneumoniae

US-09-489-039A-172

Alignment Scores:

Pred. No.: 25 Length: 825
Score: 47.00 Matches: 9
Percent Similarity: 91.67% Conservative: 2
Best Local Similarity: 75.00% Mismatches: 1
Query Match: 67.14% Indels: 0
DB: 4 Gaps: 0

US-09-017-715a-2_COPY_94_107 (1-14) x US-09-489-039A-172 (1-825)

QY 3 ArgLysGluAspLeuArgProSerAlaProGlnGln 14
|||

Db 234 CGATTGAGCATCTCGCGCCGCTGGCGCACGCCAG 199

CURRENT APPLICATION NUMBER: US/09/103,840A
CURRENT FILING DATE: 1998-06-24
NUMBER OF SEQ ID NOS: 2
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 2
LENGTH: 4403765
TYPE: DNA
ORGANISM: Mycobacterium tuberculosis
FEATURE:
OTHER INFORMATION: CDC 1551
OTHER INFORMATION: "n" bases at various positions throughout the sequence
OTHER INFORMATION: represent a, t, c or g
US-09-103-840A-2

Alignment Scores:
Pred. No.: 7.98e+05 Length: 4403765
Score: 45.00 Matches: 9
Percent Similarity: 78.57% Conservative: 2
Best Local Similarity: 64.29% Mismatches: 3
Query Match: 64.29% Indels: 0
Gaps: 0
DB: 3

US-09-017-715A-2_COPY_94_107 (1-14) x US-09-103-840A-2 (1-4403765)

Oy 1 ValValArgLyGluAspLeuArgProSerAlaProGlnGln 14
Db 3343973 GTGTTGCGTGGCGAGACCTGCTGCGCGCGCGCGCAG 3343932

RESULT 9
US-09-103-840A-1/c
Sequence 1, Application US/09103840A
Patent No. 6294328
GENERAL INFORMATION:
APPLICANT: FLEISCHMAN, Robert D.
APPLICANT: WHITE, Owen R.
APPLICANT: FRASER, Claire M.
APPLICANT: VENTER, John C.
TITLE OF INVENTION: DNA SEQUENCES FOR STRAIN ANALYSIS IN MYCOBACTERIUM
FILE REFERENCE: 24366-20007.00
CURRENT APPLICATION NUMBER: US/09/103,840A
CURRENT FILING DATE: 1998-06-24
NUMBER OF SEQ ID NOS: 2
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 1
LENGTH: 4411529
TYPE: DNA
ORGANISM: Mycobacterium tuberculosis
OTHER INFORMATION: H37Rv
US-09-103-840A-1

Alignment Scores:
Pred. No.: 7.99e+05 Length: 4411529
Score: 45.00 Matches: 9
Percent Similarity: 78.57% Conservative: 2
Best Local Similarity: 64.29% Mismatches: 3
Query Match: 64.29% Indels: 0
Gaps: 0
DB: 3

US-09-017-715A-2_COPY_94_107 (1-14) x US-09-103-840A-1 (1-4411529)

Oy 1 ValValArgLyGluAspLeuArgProSerAlaProGlnGln 14
Db 3349641 GTGTTGCGTGGCGAGACCTGCTGCGCGCGCGCAG 3349600

RESULT 10
US-09-902-540-8804
Sequence 8804, Application US/0902540
Patent No. 6833447
GENERAL INFORMATION:
APPLICANT: Goldman, Barry S.
APPLICANT: Hinkle, Gregory J.
APPLICANT: Slater, Steven C.

APPLICANT: Wiegand, Roger C.
TITLE OF INVENTION: Myxococcus xanthus Genome Sequences and Uses Thereof
FILE REFERENCE: 38-10(15849)B
CURRENT APPLICATION NUMBER: US/09/902,540
CURRENT FILING DATE: 2001-07-10
PRIOR APPLICATION NUMBER: 60/217,883
PRIOR FILING DATE: 2000-07-10
NUMBER OF SEQ ID NOS: 16825
SEQ ID NO 8804
LENGTH: 1011
TYPE: DNA
ORGANISM: Myxococcus xanthus
US-09-902-540-8804

Alignment Scores:
Pred. No.: 161 Length: 1011
Score: 43.00 Matches: 7
Percent Similarity: 84.62% Conservative: 4
Best Local Similarity: 53.85% Mismatches: 2
Query Match: 61.43% Indels: 0
Gaps: 0
DB: 4

US-09-017-715A-2_COPY_94_107 (1-14) x US-09-902-540-8804 (1-1011)

Oy 2 ValArgLyGluAspLeuArgProSerAlaProGlnGln 14
Db 46 ATGCCGAGCCCGAGCGTGTGCGCGCGCGCGCGAGCGC 84

RESULT 11
US-08-440-856A-2
Sequence 2, Application US/08440856A
Patent No. 5750873
GENERAL INFORMATION:
APPLICANT: DELAPORTA, STEPHEN L.
TITLE OF INVENTION: MATERIALS AND METHODS FOR PRODUCING
NUMBER OF SEQUENCES: 9
TITLE OF INVENTION: PLANTS WITH SINGLE-SEX FLOWERS
CORRESPONDENCE ADDRESS:
ADDRESSEE: MORRISON & FOERSTER
STREET: 2000 PENNSYLVANIA AVE. N.W.
CITY: WASHINGTON
STATE: D.C.
COUNTRY: USA
ZIP: 20037
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/440,856A
FILING DATE: 15-MAY-1995
CLASSIFICATION: 800
ATTORNEY/AGENT INFORMATION:
NAME: MILLMAN, ROBERT A.
REGISTRATION NUMBER: 36,217
REFERENCE/DOCKET NUMBER: 05463-20001.00
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202) 887-1517
TELEFAX: (202) 887-0763
TELEX: 706141
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 1187 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-440-856A-2

Alignment Scores:
Pred. No.: 193 Length: 1187
Score: 43.00 Matches: 8
Percent Similarity: 78.57% Conservative: 3

Best Local Similarity: 57.14% Mismatches: 3
Query Match: 61.43% Indels: 0
DB: 1 Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x US-08-440-856A-2 (1-1187)

QY 1 ValValArgLysGluAspLeuArgProSerAlaProGlnGln 14
DB 467 GTCTTCGACCCCGGAGATTGACCGCTCTCCGCTCA 508

RESULT 12
US-09-902-540-935/C
Sequence 935, Application US/09902540
Patent No. 6833447
GENERAL INFORMATION:
APPLICANT: Goldman, Barry S.
APPLICANT: Hinkle, Gregory J.
APPLICANT: Slater, Steven C.
APPLICANT: Miesgard, Roger C.
TITLE OF INVENTION: Myxococcus xanthus Genome Sequences and Uses Thereof
FILE REFERENCE: 38-10(115849)B
CURRENT APPLICATION NUMBER: US/09/902,540
CURRENT FILING DATE: 2001-07-10
PRIOR APPLICATION NUMBER: 60/217,883
PRIOR FILING DATE: 2000-07-10
NUMBER OF SEQ ID NOS: 16825
SEQ ID NO 935
LENGTH: 10096
TYPE: DNA
ORGANISM: Myxococcus xanthus
FEATURE:
NAME/KEY: unsure
LOCATION: (1)..(10096)
OTHER INFORMATION: unsure at all n locations
US-09-902-540-935

Alignment Scores:
Pred. No.: 2.19e+03 Length: 10096
Score: 43.00 Matches: 7
Percent Similarity: 84.62% Conservative: 4
Best Local Similarity: 53.85% Mismatches: 2
Query Match: 61.43% Indels: 0
Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x US-09-902-540-935 (1-10096)

QY 2 ValArgLysGluAspLeuArgProSerAlaProGlnGln 14
DB 2820 ATGGCGAAGCCCGGACGTGCGCCCGCGCGCCCGGAGCGC 2782

RESULT 13
US-09-949-016-5443
Sequence 5443, Application US/09949016
Patent No. 6812339
GENERAL INFORMATION:
APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
FILE REFERENCE: C1001307
CURRENT APPLICATION NUMBER: US/09/949,016
CURRENT FILING DATE: 2000-04-14
PRIOR APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 60/231,498
PRIOR FILING DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 5443
LENGTH: 689
TYPE: DNA
ORGANISM: Human

US-09-949-016-5443

Alignment Scores:
Pred. No.: 156 Length: 689
Score: 42.00 Matches: 8
Percent Similarity: 90.00% Conservative: 1
Best Local Similarity: 80.00% Mismatches: 1
Query Match: 60.00% Indels: 0
Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x US-09-949-016-5443 (1-689)

QY 3 ArgLysGluAspLeuArgProSerAlaPro 12
DB 156 AGGAGAGGACGCTGCGCCCATCAGCTCC 185

RESULT 14
US-09-419-568F-24
Sequence 24, Application US/09419568F
Patent No. 6311613
GENERAL INFORMATION:
APPLICANT: Dumoutier, Laure
APPLICANT: Louhed, Jamila
APPLICANT: Renaud, Jean-Christophe
TITLE OF INVENTION: Isolated Nucleic Acid Molecules which Encode T Cell Inducible Fac
TITLE OF INVENTION: (TIFs) The Proteins Encoded, and Uses Thereof
FILE REFERENCE: LUD 5543.2
CURRENT APPLICATION NUMBER: US/09/419,568F
CURRENT FILING DATE: 1999-10-18
PRIOR APPLICATION NUMBER: US09/354,243
PRIOR FILING DATE: 1999-07-16
PRIOR APPLICATION NUMBER: US09/178,973
PRIOR FILING DATE: 1998-10-26
NUMBER OF SEQ ID NOS: 29
SEQ ID NO 24
LENGTH: 690
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
US-09-419-568F-24

Alignment Scores:
Pred. No.: 156 Length: 690
Score: 42.00 Matches: 8
Percent Similarity: 90.00% Conservative: 1
Best Local Similarity: 80.00% Mismatches: 1
Query Match: 60.00% Indels: 0
Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x US-09-419-568F-24 (1-690)

QY 3 ArgLysGluAspLeuArgProSerAlaPro 12
DB 157 AGGAGAGGACGCTGCGCCCATCAGCTCC 186

RESULT 15
US-09-354-243B-24
Sequence 24, Application US/09354243B
Patent No. 6359117
GENERAL INFORMATION:
APPLICANT: Dumoutier, Laure
APPLICANT: Louhed, Jamila
APPLICANT: Renaud, Jean-Christophe
TITLE OF INVENTION: Isolated Nucleic Acid Molecules which Encode T Cell Inducible Fac
TITLE OF INVENTION: (TIFs) The Proteins Encoded, and Uses Thereof
FILE REFERENCE: LUD 5543.1
CURRENT APPLICATION NUMBER: US/09/354,243B
CURRENT FILING DATE: 1999-07-16
PRIOR APPLICATION NUMBER: US09/178,973
PRIOR FILING DATE: 1998-10-26
NUMBER OF SEQ ID NOS: 29
SEQ ID NO 24

; LENGTH: 690
 ; TYPE: DNA
 ; ORGANISM: Homo sapiens
 ; FEATURE:
 US-09-354-243B-24

Alignment Scores:
 Pred. No.: 156 Length: 690
 Score: 42.00 Matches: 8
 Percent Similarity: 90.00% Conservative: 1
 Best Local Similarity: 80.00% Mismatches: 1
 Query Match: 60.00% Indels: 0
 Gaps: 0
 DB: 3

US-09-017-715A-2_COPY_94_107 (1-14) x US-09-354-243B-24 (1-690)

QY 3 ArgLysGluAspLeuArgProSerAlaPro 12
 |||::||| |||||
 DB 157 AGGAGAGAGAGAGCTGCGCCCATCAGCTCCC 186

Search completed: May 4, 2005, 09:42:48
 Job time : 667.416 secs

GenCore version 5.1.6
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OM protein - nucleic search, using frame_pjn model

Run on: May 4, 2005, 09:07:52 ; Search time 102.322 Seconds

(without alignments)
809.955 Million cell updates/sec

Title: US-09-017-715A-2_COPY_94_107

Perfect score: 70

Sequence: 1 VVRKEDLRSPAPQ 14

Scoring table: BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 45 summaries

Command line parameters:

-MODEL=frame+pn.model -DEV=x1h
-Q=/c912_1/USPTO.spool.h/US9017715/runat_04052005_100743_25600/app_query.fasta.1.661
-DB=N Geneseq -QFMT=fastcap -SUFFIX=ring -MINMATCH=0.1 -LOOPL=0 -LOOPEXT=0
-UNITS=bits -START=1 -END=1 -MATRIX=blom62 -TRANS=human40.cdi -LIST=45
-DOALIGN=200 -THR SCORE=ppct -THR MAX=100 -THR MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTFMT=pcio -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000
-USRR=US09017715_0CGN_1_1_703_0runat_04052005_100743_25600 -NCPU=6 -ICPU=3
-NO MMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSPELOCK=100 -LONGLOG
-DEV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

N_Geneseq_16Dec04: *
1: geneseqn1980s: *
2: geneseqn1990s: *
3: geneseqn2000s: *
4: geneseqn2001as: *
5: geneseqn2001bs: *
6: geneseqn2002as: *
7: geneseqn2002bs: *
8: geneseqn2003as: *
9: geneseqn2003bs: *
10: geneseqn2003cs: *
11: geneseqn2003ds: *
12: geneseqn2004as: *
13: geneseqn2004bs: *

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	70	100.0	479	9 ACH15493	ACH15493 Human adu
2	70	100.0	488	12 ADM66887	Adm66887 Human hom
3	70	100.0	550	2 AAV42669	AAV42669 Human bre
4	70	100.0	550	3 AAA39470	AAA39470 Human HBG
5	70	100.0	550	6 ABL63343	ABL63343 Breast ca

6	70	100.0	550	6 ABV73813	ABV73813 Human gam
7	70	100.0	550	6 ABV73915	ABV73915 Human gam
8	70	100.0	550	10 AD63558	Ad63558 Human amy
9	70	100.0	550	10 ADG47636	Adg47636 Human amy
10	70	100.0	720	2 AAX29997	Aax29997 Human per
11	70	100.0	720	6 ABS76519	Ab876519 cDNA enco
12	70	100.0	720	10 ADR43864	Ad43864 Human SNC
13	70	100.0	720	12 ADH54342	Adh54342 Human SNC
14	70	100.0	783	4 AA193778	AA193778 Human pol
15	70	100.0	796	3 AAF21784	AAf21784 Human bre
16	70	100.0	990	13 ADR98806	Adr98806 lung spec
17	54	77.1	677	2 AAX04876	Aax04876 Human gam
18	53	75.7	478	3 AAF21785	AAf21785 Human bre
19	53	75.7	4606	6 ABT10161	ABt10161 Human bre
20	53	75.7	5666	10 ADR43467	Ad43467 Human SNC
21	53	75.7	5666	10 ADR43468	Ad43468 Polymorph
22	53	75.7	5666	12 ADH53945	Adh53945 Human SNC
23	53	75.7	5666	12 ADH53946	Adh53946 Human IDE
24	53	75.7	6012	10 ADR43314	Ad43314 Human SNC
25	53	75.7	6012	12 ADH54356	Adh54356 Human SNC
26	49	70.0	787	2 AAT51183	AAt51183 Human bre
27	48	68.6	135638	10 ABX34289	ABx34289 S. atrool
28	47	67.1	657	8 ACA35809	ACA35809 Prokaryot
29	47	67.1	727	2 AAX29998	Aax29998 Mouse per
30	47	67.1	727	12 ADM66886	Adm66886 Murine ad
31	47	67.1	825	11 ACH94377	ACH94377 Klebsell
32	45	64.3	429	8 ACA01712	ACA01712 C. glutam
33	45	64.3	465	4 AA116026	AA116026 Probe #59
34	45	64.3	465	4 ABA58473	ABa58473 Human foe
35	45	64.3	465	4 AA138127	AA138127 Probe #68
36	45	64.3	465	4 ABA27546	ABa27546 Probe #60
37	45	64.3	465	4 AAK32274	AAk32274 Human bon
38	45	64.3	465	4 AAK06585	AAk06585 Human bra
39	45	64.3	465	4 ABS31980	ABs31980 Human liv
40	45	64.3	465	6 ABS07052	ABs07052 Human gen
41	45	64.3	1410	8 ACA37685	ACA37685 Prokaryot
42	45	64.3	1435	8 ACA38395	ACA38395 Prokaryot
43	45	64.3	1473	8 ACA04765	ACA04765 Prokaryot
44	45	64.3	1488	5 AAH66390	AAh66390 C glutami
45	45	64.3	1491	8 ACA96667	ACA96667 Prokaryot

ALIGNMENTS

RESULT 1	
ACH15493	
ID	ACH15493 standard; cDNA; 479 BP.
XX	
AC	ACH15493;
XX	
DT	13-OCT-2003 (first entry)
XX	
DE	Human adult brain cDNA #2705.
XX	
KM	Human; ss: sequencing by hybridisation; SBH; expressed sequence tag; EST;
XX	genome mapping; biodiversity; genetic disorder.
OS	Homo sapiens.
XX	
PN	US2003073623-A1.
XX	
PD	17-APR-2003.
XX	
PF	30-JUL-2001; 2001US-00918995.
XX	
PR	30-JUL-2001; 2001US-00918995.
XX	
PA	(DRMA/) DRMANAC R T.
PA	(LABA/) LABAT I.
PA	(STAC/) STACHE-CRAIN B.
PA	(DICK/) DICKSON M C.
XX	(JONE/) JONES L W.
XX	

PI Drmanac RT, Labat I, Stache-Crain B, Dickeon MC, Jones LW;
XX WPI; 2003-615964/58.
XX
XX New polynucleotide sequences obtained from various cDNA libraries, useful
XX as hybridization probes, as oligomers for PCR, for chromosome and gene
XX mapping, in the recombinant production of protein, or in generating
XX antisenese DNA or RNA.
XX
XX Claim 1; SEQ ID NO 2705; 44pp; English.

XX
XX The invention relates to an isolated polynucleotide comprising any one of
XX 38043 cDNA sequences, appearing as ACH12789-ACH50831, whose sequence was
XX determined by the technique of SBH (sequencing by hybridisation). Also
XX included is a purified polypeptide comprising a sequence corresponding to
XX a reading frame of the novel polynucleotide. The nucleic acid sequences
XX are useful in diagnostics as expressed sequence tags (EST) for
XX identifying expressed genes or for physical mapping of the human genome,
XX in forensics, in assessing biodiversity, or in identifying mutations
XX responsible for genetic disorders and other traits. The nucleotide
XX sequences are also useful as hybridisation probes, as oligomers for PCR,
XX for chromosome and gene mapping, in the recombinant production of
XX protein, or in generating antisense DNA or RNA. The purified polypeptide
XX is useful for generating antibodies specific for it. The present sequence
XX is one of the 38043 isolated cDNA/EST sequences. Note: The sequence data
XX for this patent did not form part of the printed specification, but was
XX obtained in electronic format directly from USPTO at
XX seqdata.uspto.gov/sequence.html?DocID=20030073623

XX
XX Sequence 479 BP; 120 A; 118 C; 178 G; 56 T; 0 U; 7 Other;

Alignment Scores:
Pred. No.: 0.00451 Length: 479
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
Gaps: 0

DB: 9

US-09-017-715A-2_COPY_94_107 (1-14) x ACHI5493 (1-479)
Qy 1 ValValArgLyGluAspLeuArgProSerAlaProGlnGln 14
Db 293 GTGGTCCGCAAGAGACTTGAGCCATCTGCCCCCAACAG 334

RESULT 2

ADM66887
ID ADM66887 standard; DNA; 488 BP.

AC ADM66887;

DT 03-JUN-2004 (first entry)

XX Human homologue of murine adipocyte specific gamma synuclein DNA Seq 20.

XX human; adipocyte specific; gene; ds; adipose tissue; anti-obesity;

XX high mobility group I-C protein; HMG1-C; obesity; leptin; ob; diabetes;

XX adipogenesis; hypertension; cardiovascular disease; anorectic;

XX antidiabetic; hypotensive; gamma synuclein.

XX Homo sapiens.

XX WO2004011618-A2.

XX 05-FEB-2004.

XX 29-JUL-2003; 2003WO-US023684.

XX 29-JUL-2002; 2002US-0398785P.

XX 12-JUN-2003; 2003US-0478206P.

XX (HMG1-C) HMG1-C INC.

PI Chada K, Chouinard R, Ashar H, Sayed AMD;
XX WPI: 2004-143846/14.
XX P-PSDB; ADM67167.
XX
XX Identifying adipocyte specific genes, useful for treating obesity or
XX diabetes, and for identifying drug targets, by differential gene
XX expression analysis between adipose tissue or stromal vascular tissue of
XX mice of different genotypes.
XX
XX Claim 11; SEQ ID NO 20; 91pp; English.

XX
XX This invention relates to a novel method for identifying genes that are
XX over-expressed in adipose tissue and as such it provides targets for anti-
XX obesity pharmaceutical compositions. Specifically, it refers to a high
XX mobility group I-C protein (HMG1-C) that is associated with obesity and
XX is epistatic to leptin, furthermore, it refers to the ob gene where an
XX autosomal recessive trait is linked to obesity and diabetes. The present
XX invention describes performing differential gene expression analysis
XX between the white adipose tissue (WAT) or stromal vascular tissue (SVT)
XX of any two different mice selected from a group consisting of wild-type,
XX HMG1-C -/-, ob/ob, or HMG1-C -/- ob/ob genotype mice. Accordingly, using
XX this method novel nucleotides and the encoded proteins thereof were
XX identified that are adipocyte specific, and as such can be used for
XX preventing adipogenesis, diagnosing and treating diabetes, obesity,
XX hypertension and cardiovascular disease, as well as screening for
XX compounds that can modulate or prevent adipogenesis and treat diabetes or
XX obesity. These compositions exhibit anorectic, antidiabetic and
XX hypotensive activities. This polynucleotide sequence is a human homologue
XX of a murine adipocyte specific DNA sequence of the invention.

XX
XX Sequence 488 BP; 127 A; 119 C; 176 G; 66 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 0.0046 Length: 488
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x ADM66887 (1-488)
Qy 1 ValValArgLyGluAspLeuArgProSerAlaProGlnGln 14
Db 291 GTGGTCCGCAAGAGACTTGAGCCATCTGCCCCCAACAG 332

RESULT 3

AAV42669
ID AAV42669 standard; cDNA; 550 BP.

AC AAV42669;

DT 09-NOV-1998 (first entry)

XX Human breast cancer specific gene 1 (BCSG1) cDNA.

XX Breast cancer specific gene 1; BCSG1; human; metastasis; diagnosis;

XX therapy; genetic marker; ds.

XX Homo sapiens.

XX Key location/Qualifiers

XX CDS 12..395

XX FT /tag= a

XX WO9833915-A1.

XX 06-AUG-1998.

XX 03-FEB-1998; 98WO-US001804.

XX 03-FEB-1997; 97US-0037080P.

XX (HUMA-) HUMAN GENOME SCI INC.
 XX Ji H, Rosen CA;
 XX WPI; 1998-446811/38.
 DR P-PSDB; AAM63123.
 XX
 PT New isolated human breast cancer specific gene - used to develop products
 PT for the diagnosis, clinical management and treatment of breast cancer and
 PT metastases.
 XX
 PS Claim 4; Fig 1; 73pp; English.
 XX
 CC This cDNA clone corresponds to the transcript of the newly identified
 CC human breast cancer specific gene 1 (BCSG1), and includes an open reading
 CC frame for a 14.2 kDa protein (see AAM63123). It was isolated from a
 CC breast cancer cDNA library following an EST search for novel genes
 CC differentially expressed in breast cancer versus healthy breast tissue.
 CC The clone is deposited at ATCC 97175 and ATCC 97856. A gradient and stage
 CC -specific BCSG1 expression has been demonstrated from virtually no
 CC detectable expression in normal or benign breast to low level and partial
 CC expression in low grade in situ breast carcinoma and high expression in
 CC infiltrating malignant breast carcinomas. BCSG1 is useful as a breast
 CC cancer progression marker. Recombinant vectors and host cells useful for
 CC recombinant production of BCSG1 polypeptides (including epitope-bearing
 CC polypeptides) are provided. BCSG1 polynucleotides, polypeptides and
 CC antibodies can be used for the detection of breast cancer cells or breast
 CC cancer metastasis, and to develop methods for the clinical management and
 CC treatment of breast cancer
 XX
 SQ Sequence 550 BP; 132 A; 145 C; 192 G; 81 T; 0 U; 0 Other;

Alignment Scores:
 Pred. No.: 0.00526 Length: 550
 Score: 70.00 Matches: 14
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 2 Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x AAV42669 (1-550)

QY 1 ValValArgLysGluAspLeuArgProSerAlaProGlnGln 14
 Db 291 GTGGTGGCAAGAGGACCTTGAGGCATCTGCCCCCAACAG 332

RESULT 4

AAA39470
 ID AAA39470 standard; DNA; 550 BP.

XX AAA39470;

XX 24-AUG-2000 (first entry)

XX Human HBGBA67A DNA.

XX Human; ADA2; cytostatic; gene therapy; treatment; cancer;
 KW amyloid-like protein; ss.

XX Homo sapiens.

XX Homo sapiens.

XX Key Location/Qualifiers

XX CDS 12..395
 FT /*tag= a
 FT /product= "HBGBA67"

XX US6054289-A.

XX 25-APR-2000.

XX 30-AUG-1996; 96US-00705771.

PR 30-AUG-1995; 95US-0002993P.
 XX (HUMA-) HUMAN GENOME SCI INC.
 XX Moore PA;
 XX WPI; 2000-338491/29.
 DR P-PSDB; AAY87779.
 XX
 PT New polynucleotide encoding human AD2 is useful for treating cancer and
 PT for isolating cDNAs and genes having similar biological activity.
 XX
 PS Disclosure; Col 27-28; 54pp; English.
 XX
 CC This invention describes a novel polynucleotide (I) encoding human AD2.
 CC The products of the invention have cytostatic activity and can be used
 CC for gene therapy. (I) is useful for treating cancer; as primers and
 CC probes for isolating full length cDNA and genes having similar biological
 CC activity. This sequence encodes a polypeptide derived from the human
 CC HBGBA67A clone which is an amyloid-like protein found in breast tissue
 XX
 SQ Sequence 550 BP; 132 A; 145 C; 192 G; 81 T; 0 U; 0 Other;

Alignment Scores:
 Pred. No.: 0.00526 Length: 550
 Score: 70.00 Matches: 14
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 3 Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x AAA39470 (1-550)

QY 1 ValValArgLysGluAspLeuArgProSerAlaProGlnGln 14
 Db 291 GTGGTGGCAAGAGGACCTTGAGGCATCTGCCCCCAACAG 332

RESULT 5

ABL63343
 ID ABL63343 standard; DNA; 550 BP.

XX ABL63343;

XX 15-MAY-2002 (first entry)

XX Breast cancer related gene sequence SEQ ID NO:1680.

XX Human; cancer; colon; breast; ovary; oesophagus; kidney; thyroid;
 KW stomach; lung; prostate; pancreas; carcinoma; antitumour; cancerous;
 KW cytostatic; gene therapy; antineoplastic; Wilms tumour; adenocarcinoma;
 KW gene; ds.

XX Homo sapiens.

XX Homo sapiens.

XX WO200194629-A2.

XX 13-DEC-2001.

XX 30-MAY-2001; 2001WO-US010838.

XX 05-JUN-2000; 2000US-0209473P.

XX 18-SEP-2000; 2000US-0209531P.

XX 18-SEP-2000; 2000US-0233133P.

XX 20-SEP-2000; 2000US-0233617P.

XX 20-SEP-2000; 2000US-0234009P.

XX 20-SEP-2000; 2000US-0234034P.

XX 20-SEP-2000; 2000US-0234052P.

XX 22-SEP-2000; 2000US-0234509P.

XX 22-SEP-2000; 2000US-0234567P.

XX 25-SEP-2000; 2000US-0234923P.

XX 25-SEP-2000; 2000US-0235077P.

XX 25-SEP-2000; 2000US-0235082P.

PR 25-SEP-2000; 2000US-0235134P.
 PR 25-SEP-2000; 2000US-0235280P.
 PR 26-SEP-2000; 2000US-0235637P.
 PR 26-SEP-2000; 2000US-0235638P.
 PR 27-SEP-2000; 2000US-0235711P.
 PR 27-SEP-2000; 2000US-0235720P.
 PR 27-SEP-2000; 2000US-0235840P.
 PR 27-SEP-2000; 2000US-0235863P.
 PR 28-SEP-2000; 2000US-0236028P.
 PR 28-SEP-2000; 2000US-0236032P.
 PR 28-SEP-2000; 2000US-0236033P.
 PR 28-SEP-2000; 2000US-0236034P.
 PR 28-SEP-2000; 2000US-0236109P.
 PR 28-SEP-2000; 2000US-0236111P.
 PR 29-SEP-2000; 2000US-0236642P.
 PR 29-SEP-2000; 2000US-0236891P.
 PR 02-OCT-2000; 2000US-0237172P.
 PR 02-OCT-2000; 2000US-0237173P.
 PR 02-OCT-2000; 2000US-0237278P.
 PR 02-OCT-2000; 2000US-0237294P.
 PR 02-OCT-2000; 2000US-0237295P.
 PR 02-OCT-2000; 2000US-0237316P.
 PR 03-OCT-2000; 2000US-0237425P.
 PR 03-OCT-2000; 2000US-0237598P.
 PR 03-OCT-2000; 2000US-0237604P.
 PR 03-OCT-2000; 2000US-0237606P.
 PR 03-OCT-2000; 2000US-0237608P.
 PR 01-NOV-2000; 2000US-0244867P.
 PR 01-NOV-2000; 2000US-0245084P.
 PA (AVAL-) AVALON PHARM.
 PI Young PE, Augustus M, Carter KC, Edner R, Endress G, Horrigan S;
 PI Sopet DR, Weaver Z;
 XX WPI; 2002-188264/24.
 DR
 XX
 PT Screening for anti-neoplastic agent involves exposing cells to a chemical
 PT agent to be tested for anti-neoplastic activity, and determining a change
 PT in expression of a gene of a signature gene set.

Claim 1; SEQ ID NO 1680; 44pp; English.

CC The present invention describes a method (M1) for screening for an anti-
 CC neoplastic agent. The method involves exposing cells to a chemical agent
 CC to be tested for anti-neoplastic activity, determining a change in (I)
 CC expression of at least one gene (I) of a signature gene set, where (I)
 CC comprises a sequence (S) selected from 8447 sequences (given in ABL6164
 CC to ABL70110), or is at least 95% identical to (S), where a change in
 CC expression is indicative of anti-neoplastic activity. (I) has cytostatic
 CC activity and can be used in gene therapy. M1 can be used for screening an
 CC anti-neoplastic agent, and can be used for producing a product which is
 CC the data collected with respect to the anti-neoplastic agent as a result
 CC of M1, and the data is sufficient to convey the chemical structure and/or
 CC properties of the agent. M1 can be used in the treatment of cancer such
 CC as colon, breast, stomach, lung, thyroid, oesophageal, ovarian, kidney,
 CC prostate or pancreatic cancer, adenocarcinoma, carcinoma, clear cell
 CC cancer, infiltrating ductal cancer, infiltrating lobular cancer, squamous
 CC cell carcinoma, neuroendocrine carcinoma, papillary carcinoma and Wilms
 CC tumour
 XX

Sequence 550 BP; 132 A; 145 C; 192 G; 81 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.: 0.00526 Length: 550
 Score: 70.00 Matches: 14
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 Db: 6 Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x ABL63343 (1-550)

5

QY 1 ValValArgLyGluAspLeuArgProSerAlaProGlnGln 14
 DB |||||
 291 GTCGTCCGACAGGAGACTTAGCGCCATCTGCCCCCAACAG 332
 RESULT 6
 ID ABL73813
 XX ABL73813 standard; cDNA; 550 BP.
 AC
 XX ABL73813;
 DT 08-JAN-2003 (first entry)
 XX
 DE Human gamma-synuclein Glu110 variant gene.
 XX
 KW Gamma-synuclein; human; single nucleotide polymorphism; SNP;
 KW schizophrenia; neuroleptic; gene; ss.
 XX
 OS Homo sapiens.
 XX
 FH Location/Qualifiers
 FT CDS
 FT 12..395
 FT /*tag= a
 FT /product= "Gamma-synuclein"
 FT /transl_except= (pos:213..215,aa:Glu)
 FT replace(340,T)
 FT /*tag= b
 FT /standard_name= "single nucleotide polymorphism"
 XX
 PN WO200275317-A2.
 XX
 PD 26-SEP-2002.
 XX
 PF 14-MAR-2002; 2002W0-EP002872.
 XX
 PR 15-MAR-2001; 2001US-0276306P.
 XX
 PA (NOVS) NOVARTIS AG.
 PA (NOVS) NOVARTIS-ERFINDUNGEN VERW GES MBH.
 PA (UYMA) UNIV MARYLAND BALTIMORE.
 XX
 PI Roberts RC, Van Oostrum J, Voshol J, Tamminga CA;
 PI WPI; 2002-750574/81.
 DR P-PSDB; ABP54932.
 DR
 XX
 PT Screening for compounds for treating or interfering with the onset of
 PT Schizophrenia Spectrum Disorders, by detecting interactions of candidate
 PT compounds with the gamma-synuclein polypeptide.
 XX
 PS Disclosure; Fig 1; 32pp; English.
 XX
 CC The present sequence is that of cDNA encoding the Glu-110 isoform of
 CC human gamma-synuclein. The invention relates to an isoform of gamma-
 CC synuclein that is caused by an A/T single nucleotide polymorphism (SNP)
 CC at position 329 of the gamma-synuclein coding sequence. This SNP causes a
 CC glutamic acid to valine change at amino acid position 110 of gamma-
 CC synuclein, and is associated with an increased susceptibility of
 CC individuals to schizophrenia spectrum disorders (SSDs). This is the first
 CC time that a genetic component of SSDs has been identified, and provides a
 CC potential target for diagnosis and treatment of schizophrenia. Gamma-
 CC synuclein polypeptides, especially those containing the E110V mutation,
 CC are used in a claimed method of screening for compounds useful for the
 CC treatment of SSDs, and gamma-synuclein expressing cells are used in a
 CC claimed method of screening for agonist or antagonist compounds. An
 CC oligonucleotide complementary to part of the gamma-synuclein coding
 CC sequence is used for the discrimination of an SNP at position 329 of the
 CC coding sequence. Gamma-synuclein polypeptides or polynucleotides are also
 CC useful for the diagnosis of SSDs, or susceptibility to SSDs, e.g. by PCR
 CC amplification of a polynucleotide encoding gamma-synuclein and analysis
 CC of the occurrence of the SNP at position 329. A transgenic animal useful
 CC for the study of SSDs is also claimed
 XX
 SQ Sequence 550 BP; 132 A; 145 C; 192 G; 81 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.: 0.00526 Length: 550
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-09-017-715a-2_COPY_94_107 (1-14) x ABV73915 (1-550)

Qy 1 ValValArghySGluApleuArgProserAlaProGlnGln 14
Db 291 GTGTCGCAAGAGAGACTTGAGCCATCTGCCCCCAACAG 332

RESULT 7
ABV73915
ID ABV73915 standard; cDNA; 550 BP.

XX AC ABV73915;

XX DT 08-JAN-2003 (first entry)

XX DE Human gamma-synuclein Val110 variant gene.

XX KM Gamma-synuclein; human; single nucleotide polymorphism; SNP;
XX KW schizophrenia; neuroleptic; mutant; gene; ss.

XX OS Homo sapiens.

XX FH Key Location/Qualifiers

XX FT CDS 12..395
/*tag= a
/product= "Gamma-synuclein"

XX FT Variation /transl_except= (pos:213..215,aa:Glu)
/replace(340,A)

XX FT /*tag= b
/standard_name= "Single nucleotide polymorphism"

XX PN WO200275317-A2.

XX PD 26-SEP-2002.

XX PF 14-MAR-2002; 2002MO-EP002872.

XX PR 15-MAR-2001; 2001US-0276306P.

XX PA (NOVS) NOVARTIS AG.
XX PA (NOVS) NOVARTIS-ERFINDUNGEN VERW GBS MBH.
XX PA (UYMA-) UNIV MARYLAND BALTIMORE.

XX PI Roberts RC, Van Oostrum J, Voshol J, Tamminga CA;

XX DR WPI; 2002-750574/81.

XX DR P-PSDB; ABP54933.

XX PT Screening for compounds for treating or interfering with the onset of
XX PT Schizophrenia Spectrum Disorders, by detecting interactions of candidate
XX PT compounds with the gamma-synuclein polypeptide.

XX PS Disclosure; Page; 32pp; English.

XX CC The present sequence is that of cDNA encoding the Val-110 isoform of
XX CC human gamma-synuclein. The invention relates to an isoform of gamma-
XX CC synuclein that is caused by an A/T single nucleotide polymorphism (SNP)
XX CC at position 329 of the gamma-synuclein coding sequence. This SNP causes a
XX CC glutamic acid to valine change at amino acid position 110 of gamma-
XX CC synuclein, and is associated with an increased susceptibility of
XX CC individuals to schizophrenia spectrum disorders (SSDs). This is the first
XX CC time that a genetic component of SSDs has been identified, and provides a
XX CC potential target for diagnosis and treatment of schizophrenia. Gamma-
XX CC synuclein polypeptides, especially those containing the E110V mutation,
XX CC are used in a claimed method of screening for compounds useful for the

CC treatment of SSDs, and gamma-synuclein expressing cells are used in a
CC claimed method of screening for agonist or antagonist compounds. An
CC oligonucleotide complementary to part of the gamma-synuclein coding
CC sequence is used for the discrimination of an SNP at position 329 of the
CC coding sequence. Gamma-synuclein polypeptides or polynucleotides are also
CC useful for the diagnosis of SSDs, or susceptibility to SSDs, e.g. by PCR
CC amplification of a polynucleotide encoding gamma-synuclein and analysis
CC of the occurrence of the SNP at position 329. A transgenic animal useful
CC for the study of SSDs is also claimed. Note: The present sequence is not
CC shown in the specification but is derived from the gamma-synuclein
XX sequence given in Fig 1 (see ABV73915)

XX SQ Sequence 550 BP; 131 A; 145 C; 192 G; 82 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 0.00526 Length: 550
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-09-017-715a-2_COPY_94_107 (1-14) x ABV73915 (1-550)

Qy 1 ValValArghySGluApleuArgProserAlaProGlnGln 14
Db 291 GTGTCGCAAGAGAGACTTGAGCCATCTGCCCCCAACAG 332

RESULT 8
AAD63568
ID AAD63568 standard; cDNA; 550 BP.

XX AC AAD63568;

XX DT 12-FEB-2004 (first entry)

XX DE Human amyloid-like protein cDNA.

XX KM Human; genetic disease; muscular dystrophy; cystic fibrosis; cytostatic;
XX KW scientific research; gene therapy; gene; amyloid-like protein; ss.

XX OS Homo sapiens.

XX FH Key Location/Qualifiers

XX FT CDS 12..395
/*tag= a
/product= "Human amyloid-like protein"

XX PN US6639052-B1.

XX PD 28-OCT-2003.

XX PF 14-OCT-1999; 99US-00417540.

XX PR 30-AUG-1995; 95US-0002993P.
XX PR 30-AUG-1996; 96US-00705771.

XX PA (HUMA-) HUMAN GENOME SCI INC.

XX PI Moore PA;

XX DR WPI; 2003-842790/78.

XX DR P-PSDB; ABW02024.

XX PT New isolated protein and nucleic acid molecules, useful for diagnostic
XX PT and therapeutic purposes, e.g. for treating genetic diseases such as
XX PT muscular dystrophy or cystic fibrosis.

XX PS Example 5; Fig 1; Opp; English.

XX CC The invention relates to isolated new isolated protein and nucleic acid
XX CC molecules useful for diagnostic and therapeutic purposes. The invention
XX CC is for treating genetic diseases such as muscular dystrophy or cystic

CC fibrosis, and for in vitro purposes related to scientific research,
CC synthesis of DNA and manufacture of DNA vectors. The invention is useful
CC in gene therapy. The present sequence is human amyloid-like protein cDNA

XX SQ Sequence 550 BP; 132 A; 145 C; 192 G; 81 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.:	0.00526	Length:	550
Score:	70.00	Matches:	14
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	10	Gaps:	0

US-09-017-715a-2_COPY_94_107 (1-14) x AAD63568 (1-550)

OY 1 ValValArgLySGluAspLeuArgProSerAlaProGlnGln 14
DB 291 GTGGTCCGACAGAGACTTGAGCCATCTGCCCCCAACAG 332

RESULT 9

ADG47636
ID ADG47636 standard; cDNA; 550 BP.

AC ADG47636;

DT 11-MAR-2004 (first entry)

DE Human amyloid like protein cDNA.

XX ss; gene; muscular dystrophy; cystic fibrosis; hypertension;
XX angina pectoris; myocardial infarction; ulcer; asthma; allergy;
XX psychos; depression; migraine; vomiting; benign prostatic hypertrophy;
XX osteoporosis; human.

OS Homo sapiens.

XX Key Location/Qualifiers

XX FT CDS 12..395
XX FT /tag= a
XX FT /product= "Amyloid like protein"

XX US2003208043-A1.

XX 06-NOV-2003.

XX 04-JUN-2003; 2003US-00453478.

XX 30-AUG-1995; 95US-0002993P.

XX 30-AUG-1996; 96US-00705771.

XX 14-OCT-1999; 99US-00417540.

XX (HUMA-) HUMAN GENOME SCI INC.

XX Moore PA, Gentz RL, Ji H, Ni J, Hu J;

XX MPI; 2003-664796/80.

XX P-PSDB; ADG47647.

XX New human polypeptides and polynucleotides, useful for diagnosing or
XX treating genetic diseases such as muscular dystrophy or cystic fibrosis,
XX hypertension, asthma, depression or osteoporosis.

XX Claim 18; SEQ ID NO 1; 56pp; English.

XX The invention relates to an isolated human polypeptide. The polypeptides,
XX polynucleotides, agonists or antagonist are useful for diagnosing or
XX treating genetic diseases such as muscular dystrophy or cystic fibrosis,
XX hypertension, angina pectoris, myocardial infarction, ulcers, asthma,
XX allergies, psychoses, depression, migraine, vomiting, benign prostatic
XX hypertrophy or osteoporosis. The polypeptides and polynucleotides are
XX useful for in vitro purposes related to scientific research, synthesis of
XX DNA and manufacture of DNA vector. The present sequence represents cDNA

CC encoding human amyloid like protein.

XX SQ Sequence 550 BP; 132 A; 145 C; 192 G; 81 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.:	0.00526	Length:	550
Score:	70.00	Matches:	14
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	10	Gaps:	0

US-09-017-715a-2_COPY_94_107 (1-14) x ADG47636 (1-550)

OY 1 ValValArgLySGluAspLeuArgProSerAlaProGlnGln 14
DB 291 GTGGTCCGACAGAGACTTGAGCCATCTGCCCCCAACAG 332

RESULT 10

AA29997
ID AA29997 standard; DNA; 720 BP.

AC AA29997;

DT 06-JUL-1999 (first entry)

DE Human peryn gene.

XX Human; synuclein; peryn; diagnosis; neurodegenerative disorder; cancer;
XX breast; skin; intermediate filament damage; 88.

OS Homo sapiens.

XX EP908727-A1.

XX 14-APR-1999.

XX 21-SEP-1998; 98EP-00307628.

XX 19-SEP-1997; 97GB-00019879.

XX (NEUR-) NEUROBA LTD.

XX (UYSA-) UNIV ST ANDREWS.

XX MPI; 1999-217169/19.

XX P-PSDB; AAY07271.

XX New synuclein protein (persyn) and gene, useful in assays for screening,
XX diagnosing or monitoring cancer, neurodegenerative disorders or skin
XX disorders.

XX Claim 29; Page 16-17; 39pp; English.

XX This sequence represents the gene encoding a novel human synuclein family
XX member designated persyn. The sequence is useful for screening,
XX diagnosing or monitoring cancer (especially breast or skin cancer),
XX neurodegenerative disorders or skin disorders and for identifying cells
XX having intermediate filament damage

XX SQ Sequence 720 BP; 173 A; 209 C; 215 G; 123 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.:	0.00711	Length:	720
Score:	70.00	Matches:	14
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
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US-09-017-715a-2_COPY_94_107 (1-14) x AA29997 (1-720)

OY 1 ValValArgLySGluAspLeuArgProSerAlaProGlnGln 14
|||

DB 328 GTGGTGGCAAGAGAGCTTGAGGCAATCTGCCCCCAACAG 369

RESULT 11

ABST76519

ID ABST76519 standard; cDNA; 720 BP.

XX AC ABST76519;

XX 11-DEC-2002 (first entry)

DE cDNA encoding human ovarian cancer marker OV60.

KM Human; ovarian cancer; marker; cancer; familial history; brain disorder; central nervous system disorder; bacterial meningitis; viral meningitis; Alzheimer's disease; Parkinson's disease; cerebral edema; hydrocephalus; brain herniation; inflammation; encephalitis; testicular disorder; nontuberculous granulomatous orchitis; connective tissue disorder; heart disorder; ischaemic heart disease; atherosclerosis; neoplasm; histological type; carcinogenic; ovarian cancer marker; gene; ss.

XX OS Homo sapiens.

XX PN WO200271928-A2.

XX PD 19-SEP-2002.

XX PF 14-MAR-2002; 2002WO-US007826.

XX PR 14-MAR-2001; 2001US-0276025P.

XX PR 14-MAR-2001; 2001US-0276026P.

XX PR 10-AUG-2001; 2001US-0311732P.

XX PR 19-SEP-2001; 2001US-0323580P.

XX PR 26-SEP-2001; 2001US-0324967P.

XX PR 26-SEP-2001; 2001US-0325102P.

XX PR 26-SEP-2001; 2001US-0325149P.

XX (MILL-) MILLENNIUM PHARM INC.

XX PA Monahan JE, Gannavarapu M, Hoersch S, Kamatkar S, Kovatis SG, Meyer RE, Morrissey WP, Olandt PJ, Sen A, Vieby PO, Mills GB, Bat RC, Lu K, Schmandt RE, Zhao X, Glatt K;

XX PI Bat RC, Lu K, Schmandt RE, Zhao X, Glatt K;

XX DR P-PSDB; ABG96420.

XX PT WPI: 2002-723277/78.

XX PT Assessing whether a patient is afflicted with ovarian cancer, useful in the expression level of a cancer marker in a sample from a patient and from a non cancer patient.

XX PT Disclosure; Page 411; 481pp; English.

XX The present invention relates to a new method for assessing whether a patient is afflicted with ovarian cancer. The method involves comparing the expression level of a marker in a patient sample and the normal level of expression of the marker in a control non-ovarian cancer sample, where the marker is selected from 363 cancer markers described in the specification. The method of the invention is useful in diagnosing or characterizing cancer, in detecting the presence of cancer as early as possible, and the recurrence of ovarian cancer. The method may also be of particular use with patients having an enhanced risk of developing ovarian cancer (e.g. patients having a familial history of ovarian cancer). The cancer markers may be used in the management and treatment of e.g. brain and central nervous system disorders (e.g. bacterial and viral meningitis, Alzheimer's disease or Parkinson's disease), brain disorders (e.g. cerebral edema, hydrocephalus or brain herniations), inflammations (e.g. bacterial or viral meningitis or encephalitis), testicular disorders (e.g. nontuberculous granulomatous orchitis), connective tissue disorders, or heart disorders (e.g. ischemic heart disease or atherosclerosis). The compositions and methods may also be used in assessing the histological type of neoplasm associated with ovarian cancer, monitoring the progression of ovarian cancer, determining whether ovarian cancer has metastasized or is likely to metastasize,

CC selecting a composition for inhibiting ovarian cancer, assessing the CC ovarian carcinogenic potential of a compound, or inhibiting ovarian CC cancer or at risk of developing ovarian cancer. The present nucleic acid CC sequence encodes one of the ovarian cancer markers described in the CC invention

XX SO Sequence 720 BP; 173 A; 209 C; 215 G; 123 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.	Score:	Length:	Matches:	Conservative:	Mismatches:	Indels:	Gaps:
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Query Match:	100.00%						

DB: 6

US-09-017-715a-2_copy_94_107 (1-14) x ABST76519 (1-720)

Qy 1 ValValArgbysgluapleuArqProSerAlaProGlnGln 14

DB 328 GTGGTGGCAAGAGAGCTTGAGGCAATCTGCCCCCAACAG 369

RESULT 12

ADE43864

ID ADE43864 standard; cDNA; 720 BP.

XX AC ADE43864;

XX DT 29-JAN-2004 (first entry)

XX DE Human SNGC cDNA, SEQ ID 469.

XX KM Neurodegenerative disease; uPA; SNGC; IDE; KNSL1, LIPA, TNFRSF6; KM Alzheimer's disease; neuroprotective; neurotropic; gene therapy;

XX KM Chromosome 10; gene; ss.

XX OS Homo sapiens.

XX PN WO2003054143-A2.

XX PD 03-JUL-2003.

XX PF 25-OCT-2002; 2002WO-US034679.

XX PR 25-OCT-2001; 2001US-0339525P.

XX PR 08-NOV-2001; 2001US-0336929P.

XX PR 08-NOV-2001; 2001US-0338010P.

XX PR 09-NOV-2001; 2001US-0338363P.

XX PR 04-DEC-2001; 2001US-0337052P.

XX PR 28-MAR-2002; 2002US-0368919P.

XX PA (NEUR-) NEUROGENETICS INC.

XX PA (GEHO) GEN HOSPITAL CORP.

XX PI Becker KD, Velicelebi G, Elliott KJ, Wang X, Tanzi RE, Bertram L, Saunders AJ, Mullin KM, Sampson AJ, Blacker DL;

XX DR WPI: 2003-559131/52.

XX PT Determining a predisposition for or the occurrence of neurodegenerative PT disease, e.g. Alzheimer's disease by detecting in a target nucleic acid PT the presence or absence of an allelic variant of one or more polymorphic PT regions.

XX PS Claim 84; Page 740; 848pp; English.

XX The present invention relates to a method (M1) for determining a CC predisposition for or the occurrence of neurodegenerative disease in a CC subject. The method comprises detecting in a target nucleic acid obtained CC from the subject the presence or absence of an allelic variant of one or CC more polymorphic regions of one or more genes selected from uPA CC (urokinase plasminogen activator), SNGC (gamma-synuclein), IDE (insulin- CC degrading enzyme), KNSL1 (Kinesin-like protein 1), LIPA (lysosomal acid

CC lysase), and TNFRSF6 (Tumour Necrosis Factor Receptor-SF6), where the
CC presence of at least one of the allelic variant of one or more
CC polymorphic regions is indicative of a predisposition for or the
CC occurrence of neurodegenerative disease. The genes are all located on
CC chromosome 10. M1 is useful for determining a predisposition for or the
CC occurrence of, and for treating neurodegenerative disease, particularly
CC Alzheimer's disease.

XX
SQ Sequence 720 BP; 172 A; 205 C; 212 G; 120 T; 0 U; 11 Other;

Alignment Scores:

Pred. No.:	Length:	720
Score:	70.00	Matches: 14
Percent Similarity:	100.00%	Conservative: 0
Best Local Similarity:	100.00%	Mismatches: 0
Query Match:	100.00%	Indels: 0
DB:	10	Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x ADE43864 (1-720)

QY 1 ValValArgLyGluAspLeuArgProSerAlaProGlnGln 14

Db 328 GTGGTCCGCAAGAGACTTGAAGCCATCTGCCCCCAACAG 369

RESULT 13

ADH54342
ID ADH54342 standard; cDNA; 720 BP.

XX ADH54342;

XX 25-MAR-2004 (first entry)

DE Human SNCG gene cDNA sequence SeqID469.

XX human; neurodegenerative disease; urokinase plasminogen activator; uPA;
KW gamma-synuclein; SNCG; insulin degrading enzyme; IDE;
KW kinesin-like protein 1; KNSL1; lysosomal acid lipase; LIPA;
KW tumour necrosis factor receptor SF6; TNFRSF6; Alzheimer's disease; ss.
XX Homo sapiens.

XX US2003224380-A1.

XX 04-DEC-2003.

XX 25-OCT-2002; 2002US-00282174.

XX 25-OCT-2001; 2001US-0339525P.

XX 25-OCT-2001; 2001US-0348065P.

XX 02-NOV-2001; 2001US-0336983P.

XX 08-NOV-2001; 2001US-0336929P.

XX 09-NOV-2001; 2001US-0338610P.

XX 04-DEC-2001; 2001US-0337052P.

XX 28-MAR-2002; 2002US-0368919P.

XX (GENO) GEN HOSPITAL CORP.

XX Becker KD, Velicelabi G, Elliott KI, Wang X, Tanzi RE;
PI Bertram L, Saunders AJ, Mullin KM, Sampson AJ;

XX WPI; 2004-060538/06.

XX Determining a predisposition for or the occurrence of neurodegenerative
PT disease, particularly Alzheimer's disease, comprises determining the
PT presence of a polymorphism in the uPA, SNCG, IDE, KNSL1, LIPA or TNFRSF6
PT gene.

XX Claim 84; SEQ ID NO 469; 205pp; English.

CC This invention relates to a novel method of determining a predisposition
CC for or the occurrence of neurodegenerative disease comprising detecting
CC in a target nucleic acid obtained from the subject the presence of an

CC allelic variant of polymorphic regions of human genes selected from
CC urokinase plasminogen activator (uPA), gamma-synuclein (SNCG), insulin
CC degrading enzyme (IDE), kinesin-like protein 1 (KNSL1), lysosomal acid
CC lipase (LIPA) and tumour necrosis factor receptor SF6 (TNFRSF6). The
CC method is useful in determining the presence or predisposition to a
CC neurodegenerative disease, particularly Alzheimer's disease. The present
CC sequence is the cDNA sequence of the human SNCG gene which is related to
CC the invention.

XX
SQ Sequence 720 BP; 172 A; 205 C; 212 G; 120 T; 0 U; 11 Other;

Alignment Scores:

Pred. No.:	Length:	720
Score:	70.00	Matches: 14
Percent Similarity:	100.00%	Conservative: 0
Best Local Similarity:	100.00%	Mismatches: 0
Query Match:	100.00%	Indels: 0
DB:	12	Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x ADH54342 (1-720)

QY 1 ValValArgLyGluAspLeuArgProSerAlaProGlnGln 14

Db 328 GTGGTCCGCAAGAGACTTGAAGCCATCTGCCCCCAACAG 369

RESULT 14

AAI93778
ID AAI93778 standard; cDNA; 783 BP.

XX AAI93778;

XX 06-NOV-2001 (first entry)

DE Human polynucleotide SEQ ID NO 13838.

XX Human; cytokine; cell proliferation; cell differentiation; gene therapy;
KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;
KW tissue growth factor; immunomodulatory; cancer; leukaemia;
KW nervous system disorders; arthritis; inflammation; ss.
XX Homo sapiens.

XX WO200164835-A2.

XX 07-SEP-2001.

XX 26-FEB-2001; 2001WO-US004927.

XX 28-FEB-2000; 2000US-00515126.

XX 18-MAY-2000; 2000US-00577409.

XX (HYSE-) HYSEQ INC.

XX Tang YT, Liu C, Drmanac RT;

XX WPI; 2001-514838/56.

XX P-PSDB; AAO13847.

XX Isolated nucleic acids and polypeptides, useful for preventing diagnosing
PT and treating e.g. leukemia, inflammation and immune disorders.

XX

XX Claim 1; SEQ ID NO 13838; 1399pp + Sequence Listing; English.

CC The invention relates to human polynucleotides (AAI93841) and
CC the encoded proteins (AAO00010-AAO13910) that exhibit activity elating to
CC cytokine, cell proliferation or cell differentiation or which may induce
CC production of other cytokines in other cell populations. The
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or
CC peptide therapy. The polypeptides have various cytokine-like activities,
CC e.g. stem cell growth factor activity, haematopoiesis regulating
CC activity, tissue growth factor activity, immunomodulatory activity and
CC activity/inhibit activity and may be useful in the diagnosis and/or
CC treatment of cancer, leukaemia, nervous system disorders, arthritis and

CC inflammation. Note: The sequence data for this patent did not form part
CC of the printed specification, but was obtained in electronic format
CC directly from WIPO at ftp.wipo.int/pub/published_pat_sequences

XX Sequence 783 BP, 187 A; 232 C; 237 G; 127 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.:	Length:	Matches:	Conservative:	Mismatches:	Indels:	Gaps:
Score:	70.00	14	14	0	0	0
Percent Similarity:	100.00%					
Best Local Similarity:	100.00%					
Query Match:	100.00%					

US-09-017-715A-2_COPY_94_107 (1-14) x AAF21784 (1-783)

QY 1 ValValAArgbysgluapleuArqProSeRaLaProGlnGln 14
DB 393 GTGTCGCGAAGAGAGACTTGAGGCCATCTGCCCCCAACAG 434

RESULT 15

AAAF21784
ID AAF21784 standard; DNA; 796 BP.

AC AAF21784;

DT 27-MAR-2001 (first entry)

DE Human breast and ovarian cancer associated antigen gene SEQ ID 171.

XX Human; breast cancer; ovarian cancer; cytostatic; immunosuppressive;
XX neurotropic; neuroprotective; antiviral; antiallergic; hepatotropic;
XX antidiabetic; antiinflammatory; antiviral; vulnery; anticonvulsant;
XX antibacterial; antifungal; antiparasitic; cardiant; immune disorder;
XX Addison's disease; allergy; autoimmune haemolytic anaemia;
XX autoimmune thyroiditis; diabetes mellitus; Crohn's disease;
XX multiple sclerosis; rheumatoid arthritis; ulcerative colitis;
XX cardiovascular disorder; wound healing; neurological disease; ds.

OS Homo sapiens.

PN W0200055173-A1.

PD 21-SEP-2000.

PF 08-MAR-2000; 2000WO-US005881.

PR 12-MAR-1999; 99US-0124270P.

PA (HUMA-) HUMAN GENOME SCI INC.

PI Rosen CA, Ruben SM;

DR WPI; 2000-611515/58.

P-PDSB; AAB58881.

PT New human breast and ovarian cancer associated gene sequences and the
PT polypeptides encoded by these genes, useful in the prevention, treatment
PT and diagnosis of cancer, immune disorders, cardiovascular disorders and
PT neurological diseases.

PS Claim 1; Page 608; 1299pp; English.

XX Sequences AAF21614 - AAF22031 represent DNA sequences encoding human
CC proteins AAB58711 - AAB59128. The DNA and protein sequences are
CC associated with breast and ovarian cancer. Included in the invention are
CC sequences AAF22032 - AAF22040 and AAB59129 which are used in the
CC isolation and characterization of the DNA and protein sequences of the
CC invention. The breast and ovarian cancer associated DNA, protein, agonist
CC or antagonist sequences exhibit cytostatic; immunosuppressive; neurotropic;
CC neuroprotective; antiviral; antiallergic; hepatotropic; antidiabetic;
CC antiinflammatory; antiviral; vulnery; anticonvulsant; antibacterial;
CC antifungal; antiparasitic and cardiant activity. The polynucleotide and

CC protein sequences are used in the diagnosis of cancer, particularly
CC breast and ovarian cancer. The nucleic acid sequences, proteins, agonists
CC and agonists may also be used in the diagnosis, prevention and treatment
CC of immune disorders e.g. Addison's disease, allergies, autoimmune
CC haemolytic anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's
CC disease, multiple sclerosis, rheumatoid arthritis and ulcerative colitis;
CC cardiovascular disorders such as myocardial ischaemias; wound healing;
CC neurological diseases such as cerebral anoxia and epilepsy; and
CC infectious diseases

XX Sequence 796 BP, 195 A; 233 C; 240 G; 126 T; 0 U; 2 Other;

Alignment Scores:

Pred. No.:	Length:	Matches:	Conservative:	Mismatches:	Indels:	Gaps:
Score:	70.00	14	14	0	0	0
Percent Similarity:	100.00%					
Best Local Similarity:	100.00%					
Query Match:	100.00%					

US-09-017-715A-2_COPY_94_107 (1-14) x AAF21784 (1-796)

QY 1 ValValAArgbysgluapleuArqProSeRaLaProGlnGln 14

DB 388 GTGTCGCGAAGAGAGACTTGAGGCCATCTGCCCCCAACAG 429

Search completed: May 4, 2005, 09:26:19
Job time: 105.322 secs

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OM protein - nucleic search, using frame_plus_p2n model

Run on: May 4, 2005, 09:07:55 ; Search time 795.087 Seconds
(without alignment)
853.206 Million cell updates/sec

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Listing first 45 summaries

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12: gb_sy: *
13: gb_un: *
14: gb_vi: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	70	100.0	381	9	CR541790 Homo sapi
2	70	100.0	384	9	CR541788 Homo sapi
3	70	100.0	488	9	AF411524 Homo sapi
4	70	100.0	550	6	AR412236 Sequence

5	70	100.0	550	6	AX331171	AX331171 Sequence
6	70	100.0	550	6	BD022727	BD022727 Mammariy c
7	70	100.0	550	6	AF010126	AF010126 Homo sapi
8	70	100.0	704	6	CQ720882	CQ720882 Sequence
9	70	100.0	720	6	E36333	E36333 Analytical
10	70	100.0	720	6	AX004527	AX004527 Sequence
11	70	100.0	720	9	AF017255	AF017255 Homo sapi
12	70	100.0	728	11	BV177827	BV177827 segm97020
13	70	100.0	738	11	BC014098	BC014098 Homo sapi
14	53	75.7	4606	9	AF044311	AF044311 Homo sapi
15	53	75.7	5491	9	AF037207	AF037207 Homo sapi
16	53	75.7	149668	9	AC025268	AC025268 Homo sapi
17	53	75.7	175542	2	AC150062	AC150062 Gallus ga
18	53	75.7	188771	2	AC150073	AC150073 Gallus ga
19	53	75.7	190496	2	EX510354	EX510354 Homo sapi
20	49	70.0	168860	9	AL160175	AL160175 Human DNA
21	49	70.0	219205	10	AL611985	AL611985 Mouse DNA
22	48	68.6	135638	1	AF484556	AF484556 Streptomy
23	48	68.6	211075	10	AC128668	AC128668 Mus muscu
24	48	68.6	242974	2	AC103570	AC103570 Rattus no
25	47	67.1	488	10	AF518351	AF518351 Rattus no
26	47	67.1	727	6	E36334	E36334 Analytical
27	47	67.1	727	6	AX004529	AX004529 Sequence
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31	47	67.1	788	5	BC075925	BC075925 Danio rer
32	47	67.1	825	6	AR383443	AR383443 Sequence
33	47	67.1	1252	3	AK114896	AK114896 Ciona int
34	47	67.1	1646	3	AK114539	AK114539 Ciona int
35	47	67.1	191707	2	CR847971	CR847971 Rattus rex
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37	46	65.7	154356	10	AC131119	AC131119 Mus muscu
38	46	65.7	172435	8	OSJ000163	OSJ000163 Oryza sat
39	46	65.7	189771	2	AC118514	AC118514 Rattus no
40	46	65.7	246005	2	AC095834	AC095834 Rattus no
41	46	65.7	246005	1	AP000990	AP000990 Sulfolobu
42	45	64.3	465	6	CO070159	CO070159 Sequence
43	45	64.3	465	6	CO079954	CO079954 Sequence
44	45	64.3	465	6	CQ136809	CQ136809 Sequence
45	45	64.3	465	6	CQ174616	CQ174616 Sequence

ALIGNMENTS

RESULT 1	CR541790	381 bp	MRNA	linear	PRI 29-JUN-2004
LOCUS	CR541790				
DEFINITION	Homo sapiens full open reading frame cDNA clone RZP0834B0231D for gene SNG, synuclein, gamma (breast cancer-specific protein 1); complete cds, without stopcodon.				
ACCESSION	CR541790				
VERSION	CR541790.1	GI:49456536			
KEYWORDS	Full ORF shuttle clone, Gateway(TM), complete cds.				
SOURCE	Homo sapiens (human)				
ORGANISM	Homo sapiens				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.				
AUTHORS	Halleck A., Ebert L., Moundinya, M., Schick, M., Bisenstein, S., Korn, B., Zuo, D., Hu, Y. and Labaer, V.				
TITLE	Cloning of human full open reading frames in Gateway(TM) system entry vector (pDONR201)				
JOURNAL	Unpublished				
REFERENCE	2 (bases 1 to 381)				
AUTHORS	Halleck A., Ebert L., Moundinya, M., Schick, M., Bisenstein, S., Korn, B., Zuo, D., Hu, Y. and Labaer, V.				
TITLE	Direct Submision				
JOURNAL	Submitted (28-JUN-2004) RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH, Im Neuenheimer Feld 580, D-69120 Heidelberg, Germany				

COMMENT RZPD: RZPD0834B0231D, ORFNo 3631
www.rzpd.de/cgi-bin/products/ci.cgi?cloneid=RZPD0834B0231D RZPDLIB:
Human Full ORF Clones Gateway(TM) - RZPD (kan-resist.) RZPD LIB No.
834
www.rzpd.de/cgi-bin/products/showlib.pl.cgi/response?libNo=834
Contact: Inge Axlart
RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH,
Heubnerweg 6, D-14059 Berlin, Germany
Tel: +49 30 32639 100
Fax: +49 30 32639 111
www.rzpd.de
This clone is available from RZPD;
Contact RZPD (customer.service@rzpd.de) for further information.
Clone name at Harvard Institute of Proteomics
(www.hip.harvard.edu): F1H1310940.01L
This CDS clone is part of a collection of human full ORF clones
jointly established and verified by the Harvard Institute of
Proteomics (HIP) and RZPD.
This CDS has been inserted into pDONR201 via a BP Clonase(TM)
reaction. Additional sequence has been added in front of the start
codon: attc...AAAAA GCA GGC TCC ACC (ATG).
The last codon is followed by the 3' att site: GACCCAGCTTTCTT. att
The clone is validated by full sequence check.
Compared to the reference sequence BC014098
we did not find any amino acid exchanges.
Clone distribution: http://www.rzpd.de/products/orfclones/
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ORIGIN
Alignment Scores:
Pred. No.: 0.00392 Length: 381
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0
US-09-017-715A-2_COPY_94_107 (1-14) x CR541790 (1-381)
QY 1 ValValATgLyGtLnaSpLeuArGProSerAlaProGlnGln 14
|||||
DB 280 GTGTGTGCGCAGAGAGACTTGAAGCCATCTGCCCCCAACAG 321
|||||
RESULT 2
CR541788
LOCUS CR541788 384 bp mRNA linear PRI 29-JUN-2004
DEFINITION Homo sapiens full open reading frame cDNA clone RZPD0834F0930D for
gene SNCG, synuclein, gamma (breast cancer-specific protein 1);
complete cds, incl. stopcodon.
ACCESSION CR541788
VERSION CR541788.1 GI:49456532
KEYWORDS Full ORF shuttle clone, Gateway(TM), complete cds.
SOURCE Homo sapiens (human)

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE
AUTHORS 1 (bases 1 to 384)
Halleck,A., Ebert,L., Mroundinya,M., Schick,M., Eisenstein,S.,
Neubert,P., Ketrang,K., Schatten,R., Shen,B., Henze,S., Mar,W.,
Korn,B., Zuo,D., Hu,Y. and Labaer,J.
Cloning of human full open reading frames in Gateway(TM) system
entry vector (pDONR201)
TITLE Unpublished
2 (bases 1 to 384)
Halleck,A., Ebert,L., Mroundinya,M., Schick,M., Eisenstein,S.,
Neubert,P., Ketrang,K., Schatten,R., Shen,B., Henze,S., Mar,W.,
Korn,B., Zuo,D., Hu,Y. and Labaer,J.
Direct Submission
JOURNAL Submitted (28-JUN-2004) RZPD Deutsches Ressourcenzentrum fuer
Genomforschung GmbH, Im Neuenheimer Feld 580, D-69120 Heidelberg,
Germany
COMMENT RZPD: RZPD0834F0930D, ORFNo 3605
www.rzpd.de/cgi-bin/products/ci.cgi?cloneid=RZPD0834F0930D RZPDLIB:
Human Full ORF Clones Gateway(TM) - RZPD (kan-resist.) RZPD LIB No.
834
www.rzpd.de/cgi-bin/products/showlib.pl.cgi/response?libNo=834
Contact: Inge Axlart
RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH,
Heubnerweg 6, D-14059 Berlin, Germany
Tel: +49 30 32639 100
Fax: +49 30 32639 111
www.rzpd.de
This clone is available from RZPD;
Contact RZPD (customer.service@rzpd.de) for further information.
Clone name at Harvard Institute of Proteomics
(www.hip.harvard.edu): F1H131021.01X
This CDS clone is part of a collection of human full ORF clones
jointly established and verified by the Harvard Institute of
Proteomics (HIP) and RZPD.
This CDS has been inserted into pDONR201 via a BP Clonase(TM)
reaction. Additional sequence has been added in front of the start
codon: attc...AAAAA GCA GGC TCC ACC (ATG).
The stopcodon is followed by the 3' att site: GACCCAGCTTTCTT. att
The clone is validated by full sequence check.
Compared to the reference sequence BC014098
we did not find any amino acid exchanges.
Clone distribution: http://www.rzpd.de/products/orfclones/
Location/Qualifiers
1..384
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="RZPD0834F0930D"
/clone_lib="Human Full ORF Clones Gateway(TM) - RZPD"
/lab_host="DH5Alpha"
/note="Vector: pDONR201, Site_1: attP1, Site_2: attP2"
1..384
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1..384
/gene="SNCG"
/codon_start=1
/protein_id="CAG46587.1"
/db_xref="GI:49456533"
/translation="MDVFKGFSIAKEGVGAVEKTKQGVTEAEKTEKGVVYGAKT
KENVQSVTSVAEKTEKQANAVSEAVSVNTVAKTVEAEINIAVTSGVAKEDLR
SAPQDEGASKEKEVEAEASQSGD"
ORIGIN
Alignment Scores:
Pred. No.: 0.00395 Length: 384
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0

PF 03-FEB-1998 JP 1996515053
PR 03-FEB-1997 US 60/037080
PI HONGJUN JI, CRAIG A ROSEN
PC
C12N15/09, C07K14/47, C07K16/18, C12N1/15, C12N1/19, C12N1/21, C12N5/ PC
10,
PC C12P21/02, C12Q1/68, G01N33/574, C12N15/00, C12N5/00 CC
Strandedness: Double;
CC Topology: Both;
FH Key Location/Qualifiers
FT CDS 12..392.
Location/Qualifiers
1..550
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"

ORIGIN
Alignment Scores:
Pred. No.: 0.00543 Length: 550
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x BD022727 (1-550)

QY 1 ValValArgLysGluAspLeuArgProSerAlaProGlnGln 14
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Db 291 GTGGTGGCAGAGGAGCTTGAGGCCATCTGCCCCCAACAG 332

RESULT 7
AF010126 550 bp mRNA linear PRI 26-JUL-1997
LOCUS Homo sapiens breast cancer-specific protein 1 (BCSG1) mRNA,
DEFINITION complete cds.
ACCESSION AF010126
VERSION AF010126.1 GI:2281473
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE
AUTHORS Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Bukaryota; Euteheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 550)
Ji, H., Liu, Y.E., Jia, T., Wang, M., Liu, J., Xiao, G., Joseph, B.K.,
Rosen, C., and Shi, Y.E.
Identification of a breast cancer-specific gene, BCSG1, by direct
differential cDNA sequencing
Cancer Res. 57 (4), 759-764 (1997)
9718957
9044857
2 (bases 1 to 550)
Ji, H., Liu, Y.E., Jia, T., Wang, M., Liu, J., Xiao, G., Joseph, B.K.,
Rosen, C., and Shi, Y.E.
Direct Submision
Submitted (23-JUN-1997) Ped. Res., Long Island Jewish Medical
Center, 270-05 76th Ave., New Hyde Park, NY 11040, USA
Location/Qualifiers
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/mol_type="mRNA"
/db_xref="taxon:9606"
/rname="type="breast cancer"
/note="CDNA highly abundant in a breast cancer library but
not in normal library"
1..550
/gene="BCSG1"
12..395
/gene="BCSG1"
/note="breast cancer-specific protein 1; synuclein-like;
AD amyloid-like"
/codon_start=1

/product="BCSG1 protein"
/protein_id="BAB64109.1"
/db_xref="GI:2281474"
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SAPOQEGASKEKEVEAEAOQSGD"

ORIGIN
Alignment Scores:
Pred. No.: 0.00543 Length: 550
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x AF010126 (1-550)

QY 1 ValValArgLysGluAspLeuArgProSerAlaProGlnGln 14
|||||
Db 291 GTGGTGGCAGAGGAGCTTGAGGCCATCTGCCCCCAACAG 332

RESULT 8
CQ720882 704 bp DNA linear PAT 03-FEB-2004
LOCUS Sequence 6816 from Patent WO02068579.
DEFINITION CQ720882
ACCESSION CQ720882
VERSION CQ720882.1 GI:42281739
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE
AUTHORS Venter, C.J., Adams, M.C., Li, P.W. and Myers, E.W.
TITLE Kites, such as nucleic acid arrays, comprising a majority of
humanexons or transcripts, for detecting expression and other uses
thereof
Patent: WO 02068579-A 6816 06-SEP-2002;
PE Corporation (NY) (US)
Location/Qualifiers
1..704
/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"

ORIGIN
Alignment Scores:
Pred. No.: 0.00676 Length: 704
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x CQ720882 (1-704)

QY 1 ValValArgLysGluAspLeuArgProSerAlaProGlnGln 14
|||||
Db 327 GTGGTGGCAGAGGAGCTTGAGGCCATCTGCCCCCAACAG 368

RESULT 9
E36333 720 bp DNA linear PAT 18-JUN-2001
LOCUS Analytical matter based on synuclein and novel synuclein protein.
DEFINITION E36333
ACCESSION E36333.1 GI:13022626
VERSION E36333.1 JP 1999239488-A/1.
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE
AUTHORS Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Bukaryota; Euteheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 720)

AUTHORS Andrew, S.M., Valdimia, R.B. and Arun, M.D.
TITLE Analytical matter based on synuclein and novel synuclein protein
JOURNAL Patent: JP 199239488-A 1 07-SEP-1999;
COMMENT THE UNIVERSITY COURT OF THE UNIVERSITY OF ST ANDREWS, NYUROBA LTD
OS Homo sapiens (human)
PN JP 199239488-A/1
PD 07-SEP-1999
PF 21-SEP-1998 JP 1998306283
PR 19-SEP-1997 GB 9719879.0
PI ANDREW SMITH, MAKKARION, VALDIMIA, RUVOVICH, BUCHMAN, PI ARUN
MILWARD DAVIS
PC C12N15/09, A01K67/027, C12Q1/68, G01N33/53, C12N15/00 CC
FH Key Location/Qualifiers
FT CDS (49) (432).
LOCATION/Qualifiers
1. 720
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
ORIGIN
Alignment Scores:
Pred. No.: 0.0069 Length: 720
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0
US-09-017-715a-2_COPY_94_107 (1-14) x E36333 (1-720)
Qy 1 ValValArglysgluaspleuAtrgProSerAlaProGlnGln 14
Db 328 GTGGTGGCAAGAGGAGCTTGAGGCCATCTGCCCAACAG 369
RESULT 10
LOCUS AX004527 720 bp DNA linear PAT 24-AUG-2000
DEFINITION Sequence 1 from Patent EP0908727.
ACCESSION AX004527
VERSION AX004527.1 GI:9927977
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
AUTHORS 1
TITLE Synuclein-based assay and synuclein protein
JOURNAL Patent: EP 0908727-A 1 14-APR-1999;
NEUROBA LIMITED (GB); UNIV COURT OF THE UNIVERSITY O (GB)
FEATURES
source
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/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"
49. 432
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SAPOEGVASKEKEVEAEASGSD"
ORIGIN
Alignment Scores:
Pred. No.: 0.0069 Length: 720
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-09-017-715a-2_COPY_94_107 (1-14) x AX004527 (1-720)
Qy 1 ValValArglysgluaspleuAtrgProSerAlaProGlnGln 14
Db 328 GTGGTGGCAAGAGGAGCTTGAGGCCATCTGCCCAACAG 369
RESULT 11
LOCUS AF017256 720 bp mRNA linear PRI 23-SEP-1998
DEFINITION Homo sapiens perSyn mRNA, complete cds.
ACCESSION AF017256
VERSION AF017256.1 GI:3642774
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
AUTHORS Ninkina, N.N., Alimova-Koest, M.V., Paterson, J.W., Delaney, L.,
Cohen, B.B., Imreh, S., Gnuchev, N.V., Davies, A.M., and Buchman, V.L.
TITLE Organization, expression and polymorphism of the human perSyn gene
JOURNAL Hum. Mol. Genet. 7 (9), 1417-1424 (1998)
MEDLINE 98367030
PubMed 9700196
REFERENCE 2 (bases 1 to 720)
AUTHORS Buchman, V.L.
TITLE Direct Submission
JOURNAL Submitted (04-AUG-1997) School of Biomedical Sciences, Univ. of St.
Andrews, Bute Medical Buildings, St. Andrews, Fife KY16 9TS,
Scotland
FEATURES
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49. 432
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433. 720
684. 689
706
3' UTR
polyA_signal
polyA_site
ORIGIN
Alignment Scores:
Pred. No.: 0.0069 Length: 720
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0
US-09-017-715a-2_COPY_94_107 (1-14) x AF017256 (1-720)
Qy 1 ValValArglysgluaspleuAtrgProSerAlaProGlnGln 14
Db 328 GTGGTGGCAAGAGGAGCTTGAGGCCATCTGCCCAACAG 369
RESULT 12
LOCUS BV177827 738 bp DNA linear STS 10-JUN-2004
DEFINITION sqm7020 Human DNA (Sequencem) Homo sapiens STS genomic. sequence
tagged site.
ACCESSION BV177827
VERSION BV177827.1 GI:48014020

KEYWORDS STS.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Homindae; Homo.
TITLE 1 (bases 1 to 738)
Nelson, R.M., Marnellos, G., Kammerer, S., Hoyal, C.R., Shi, M.M.,
Cantor, C.R., and Braun, A.
Large-scale validation of single nucleotide polymorphisms in Gene
Regions
Genome Res. (2004) In press
JOURNAL COMMENT
Contact: Andreas Braun
Pharmaceuticals division
Sequenom, Inc.
3595 John Hopkins Court, San Diego, CA 92121, USA
Tel: 18582029018
Fax: 18582029020
Email: abraun@sequenom.com
Primer A: No primer sequence submitted
Primer B: No primer sequence submitted
STS size: 738.

FEATURES
source
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ORIGIN
STS
Alignment Scores:
Pred. No.: 0.00705 Length: 738
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 11 Gaps: 0

US-09-017-715A-2_COPY_94_107 (1-14) x BVL77827 (1-738)

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Db 330 GTGGTGGCAGAGGACGACCTTGAGCCCATCTGCCCAACAG 371

RESULT 13
BC014098
LOCUS BC014098 758 bp mRNA linear PRI 29-JUN-2004
DEFINITION Homo sapiens synuclein, gamma (breast cancer-specific protein 1),
ACCESSION mRNA (CDNA clone MGC:20132 IMAGE:454644), complete cds.
VERSION BC014098
KEYWORDS MGC.
SOURCE BC014098.2 GI:33878363
ORGANISM Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homindae; Homo.
1 (bases 1 to 758)
Straussberg, R.L., Feingold, E.A., Gronow, L.H., Derge, J.G.,
Klausner, R.D., Collins, F.S., Wagner, L., Shennan, C.M., Schuler, G.D.,
Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K.,
Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, F.,
Datchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L.,
Stapleton, M., Soares, M.B., Donald, M.F., Casavant, T.L.,
Scheetz, T.E., Brownstein, M.U., Usdin, T.B., Toshiyuki, S.,
Carninci, P., Prange, C., Raha, S.S., Loguercio, N.A., Peters, G.J.,
Abrams, R.D., Mullany, S.J., Bosak, S.A., McEwan, P.J.,
McKernan, K.J., Malek, J.A., Gunaratne, P.H., Richards, S.,
Morley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hulyk, S.W.,
Villalón, D.K., Muzny, D.M., Sodergren, E.J., Lu, X., Gibbs, R.A.,
Fahy, J., Hellon, E., Kettelman, W., Madan, A., Rodigues, S.,
Sanchez, A., Whiting, M., Madan, A., Young, A.C., Shevchenko, Y.,
Bouffard, G.G., Blakesley, R.W., Touchman, D.W., Green, E.D.,

TITLE
JOURNAL
PUBMED
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
REMARK
Contact: MGC help desk
On Aug 19, 2003 this sequence version replaced gi:15559464.
Contact: MGC help desk
Email: cgapbs-remail.nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: National Institutes of Health Intramural
Sequencing Center (NISC),
Gaithersburg, Maryland;
Web site: http://www.nisc.nih.gov/
Contact: nisc.mgc@nih.gov
Akhter, N., Ayele, K., Beckstrom-Sternberg, S.M., Benjamin, B.,
Blakesley, R.W., Bouffard, G.G., Bren, K., Brinkley, C., Brooks, S.,
Dieckrich, N.L., Grant, S., Guan, X., Gupta, J., Haghighi, P.,
Hansen, N., Ho, S.-L., Karlins, E., Kwong, P., Latic, P., Legaspi, R.,
Maduro, Q.L., Masiello, C., Maskeri, B., Mastriani, S.D., McCloskey, J.C.,
McDowell, J., Pearson, R., Stancitop, S., Thomas, P.J., Touchman, D.W.,
Tsurgou, C., Vogt, J.L., Walker, M.A., Wetherby, K.D., Wiggins, L.,
Young, A., Zhang, L.-H. and Green, E.D.

Clone distribution: MGC clone distribution information can be found
through the I.M.A.G.E. Consortium/LNL at: http://image.llnl.gov
Series: IRAL Plate: 28 Row: P Column: 20
This clone was selected for full length sequencing because it
passed the following selection criteria: matched mRNA gi: 4507112.

FEATURES
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/db_xref="MIM:602998"
71..454
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1)"
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/db_xref="MIM:602998"
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ORIGIN
Alignment Scores:
Pred. No.: 0.00722 Length: 758
Score: 70.00 Matches: 14
Percent Similarity: 100.00% Conservative: 0


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ACCESSION      AF037207
VERSION        AF037207.1
KEYWORDS       GI:3642902
SOURCE         Homo sapiens (human)
ORGANISM       Homo sapiens
REFERENCE      Mammalia; Eutheria; Chordata; Craniata; Vertebrata; Euarchontomi;
AUTHORS        Hominidae; Primates; Catarrhini; Hominoidea; Homo.
TITLE          1 (bases 1 to 5491)
JOURNAL        Ninkina,N.N., Alimova-Kost,M.V., Paterson,J.W., Dejaney,L.,
MEDLINE        Cohen,B.B., Imreh,S., Gnuchev,N.V., Davies,A.M. and Buchman,V.L.
PUBMED         Organization, expression and polymorphism of the human peryn gene
                Hum. Mol. Genet. 7 (9), 1417-1424 (1998)
                98367030
REFERENCE      2 (bases 1 to 5491)
AUTHORS        Buchman,V.L.
TITLE          Direct Submission
JOURNAL        Submitted (06-DEC-1997) Biomedical Sciences, University of St.
                Andrews, Bute Medical Buildings, St. Andrews, Fife KY16 9TS,
                Scotland, UK
FEATURES       Location/Qualifiers
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               4876..>5169)
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               SAPQEGVATSKKEEVABEAOSGD"
               /number=2
               1544..1585
               /number=3
               1912..2039
               /number=4
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               5019
               /note="in human peryn mRNA clone H1 deposited in GenBank
               Accession Number AF017256"
               /replace="c"
               polyA_signal   5148..5153
               polyA_site     5169
ORIGIN
polyA_signal
polyA_site
Alignment Scores:
Pred. No.:      44.9      Length:    5491
Score:          53.00     Matches:    10
Percent Similarity: 100.00% Conservative: 1
Best Local Similarity: 90.91% Mismatches: 0
Query Match:    75.71% Indels: 0
DB:             9 Gaps: 0
US-09-017-715A-2_COPY_94_107 (1-14) x AF037207 (1-5491)

```

Oy 4 LysGIuAspLeuArgProSerAlaProGInGln 14
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Db 4512 CAGGAGGACTTGAGGCGCATCTGCCCCCAACAG 4544

Search completed: May 4, 2005, 11:53:27
Job time : 801.421 secs

GenCore version 5.1.6
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OM protein - nucleic search, using frame_p2n model

Run on: May 4, 2005, 09:07:55 ; Search time 454.336 Seconds
(without alignments)
853.206 Million cell updates/sec

Title: US-09-017-715A-2_COPY_120_127

Perfect score: 41
Sequence: 1 EBAQSGSD 8

Scoring table: BLOSUM62
Xgapop 10.0, Xgapext 0.5
Ygapop 10.0, Ygapext 0.5
Fgapop 6.0, Fgapext 7.0
Delop 6.0, Delext 7.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Command line parameters:
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-Q=/cgn2.1/USPTO.spool.h/US09017715/runat_04052005_100744_25608/app.query.fasta_1.661
-DB=GenBml1 -QFMT=fastap -SUFFIX=rge -MINMATCH=0.1 -LOOPCL=0 -LOOPEXT=0
-UNITS=bits -START=1 -END=1 -MATRIX=blom62 -TRANS=human40.cdd -LIST=45
-DOCALLIGN=200 -THR SCORE=pct -THR MAX=100 -THR MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTFMT=pro -NOR=ext -HEARSIZE=500 -MINLEN=0 -MAXLEN=200000000
-USER=US09017715_QCGN_1_1_3970_@runat_04052005_100744_25608 -NCPU=6 -ICPU=3
-NO MMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG
-DEV TIMEOUT=120 -MARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

GenBml: *
1: gb_da: *
2: gb_hcg: *
3: gb_in: *
4: gb_om: *
5: gb_ov: *
6: gb_pat: *
7: gb_ph: *
8: gb_pl: *
9: gb_pr: *
10: gb_ro: *
11: gb_str: *
12: gb_sy: *
13: gb_un: *
14: gb_vl: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	41	100.0	381	9	CR541790 Homo sapi
2	41	100.0	384	9	CR541788 Homo sapi
3	41	100.0	488	3	AF411524 Homo sapi
4	41	100.0	550	6	AR412236 Sequence

5	41	100.0	550	6	AX311171	AX311171 Sequence
6	41	100.0	550	6	BD022727	BD022727 Mammariy c
7	41	100.0	550	6	AF010126	AF010126 Homo sapi
8	41	100.0	704	6	CQ720882	CQ720882 Sequence
9	41	100.0	720	6	E36333	E36333 Analytical
10	41	100.0	720	6	AX004527	AX004527 Sequence
11	41	100.0	720	9	AF017256	AF017256 Homo sapi
12	41	100.0	738	11	BV177827	BV177827 Bgmn97020
13	41	100.0	758	9	BC014098	BC014098 Homo sapi
14	41	100.0	177006	2	AC102691	AC102691 Mus muscu
15	41	100.0	255619	2	AC111772	AC111772 Rattus no
16	41	100.0	286562	2	AC137025	AC137025 Rattus no
17	41	100.0	309255	2	AC133219	AC133219 Rattus no
18	38	92.7	4887	10	AY383729	AY383729 Mus muscu
19	38	92.7	10029	1	AE010406	AE010406 Mechanopy
20	38	92.7	69346	2	AC134394	AC134394 Homo sapi
21	38	92.7	133288	9	AL357503	AL357503 Human DNA
22	38	92.7	182003	9	AL355987	AL355987 Human DNA
23	38	92.7	187534	10	AC132251	AC132251 Mus muscu
24	38	92.7	210950	10	AC129317	AC129317 Mus muscu
25	38	92.7	212717	10	AL929228	AL929228 Mouse DNA
26	38	92.7	246070	2	AC135737	AC135737 Mus muscu
27	37	90.2	488	10	AY518351	AY518351 Rattus no
28	37	90.2	732	10	RN5D5YNGE	X86789 R. norvegicu
29	37	90.2	1275	6	CQ575643	CQ575643 Sequence
30	37	90.2	3275	6	CQ575642	CQ575642 Sequence
31	37	90.2	22723	2	AC019971	AC019971 Drosophi1
32	37	90.2	89837	2	AC091822	AC091822 Homo sapi
33	37	90.2	90693	9	AC005592	AC005592 Homo sapi
34	37	90.2	96437	9	AY601819	AY601819 Homo sapi
35	37	90.2	110000	2	LMFPCR36_10	Continuation (11 o
36	37	90.2	146000	9	AP005433	AP005433 Homo sapi
37	37	90.2	156870	2	AC149695	AC149695 Bos tauru
38	37	90.2	161117	3	AC008311	AC008311 Drosophi1
39	37	90.2	162338	5	BX537341	BX537341 Zebrafish
40	37	90.2	168448	3	AC009346	AC009346 Drosophi1
41	37	90.2	191734	2	AC016560	AC016560 Homo sapi
42	37	90.2	200146	2	AC073774	AC073774 Mus muscu
43	37	90.2	209523	10	AC084386	AC084386 Mus muscu
44	37	90.2	213050	1	AL646079	AL646079 Ralstonia
45	37	90.2	219405	2	AC118096	AC118096 Rattus no

ALIGNMENTS

RESULT 1
LOCUS CR541790
DEFINITION Homo sapiens full open reading frame cDNA clone RZPO834B0231D for gene SNCG, syncleins, gamma (breast cancer-specific protein 1); complete cds, without stopcodon.
ACCESSION CR541790
VERSION CR541790.1 GI:49456536
KEYWORDS Full ORF shuttle clone, Gateway(TM), complete cds.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
1 (bases 1 to 381)
Halleck, A., Ebert, L., Mkundinya, M., Schick, M., Eisenstein, S., Neubert, P., Ketrang, K., Schatten, R., Shen, B., Henze, S., Mar, W., Korn, B., Zuo, D., Hu, Y. and Labaer, J.
Cloning of human full open reading frames in Gateway(TM) system entry vector (pDONR201)
Unpublished
2 (bases 1 to 381)
Halleck, A., Ebert, L., Mkundinya, M., Schick, M., Eisenstein, S., Neubert, P., Ketrang, K., Schatten, R., Shen, B., Henze, S., Mar, W., Korn, B., Zuo, D., Hu, Y. and Labaer, J.
Direct Submission
Submitted (28-JUN-2004) RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH, Im Neuenheimer Feld 580, D-69120 Heidelberg, Germany

COMMENT RZPD; RZPD0834B0231D, ORFNo 3631
 www.rzpd.de/cgi-bin/products/cl.cgi?cloneid=RZPD0834B0231D RZPDLIB;
 Human Full ORF Clones Gateway(TM) - RZPD (kan-resist.) RZPD LIB No.
 834
 www.rzpd.de/cgi-bin/products/showlib.pl.cgi/response?libNo=834
 www.rzpd.de/products/orfclones/
 Contact: Inge Axtelt
 RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH,
 Heubnerweg 6, D-14059 Berlin, Germany
 Tel: +49 30 32639 100
 Fax: +49 30 32639 111
 www.rzpd.de
 This clone is available from RZPD;
 Contact RZPD (customer.service@rzpd.de) for further information.
 Clone name at Harvard Institute of Proteomics
 (www.hip.harvard.edu): FLH130940.01L
 This CDS clone is part of a collection of human full ORF clones
 jointly established and verified by the Harvard Institute of
 Proteomics (HIP) and RZPD.
 This CDS has been inserted without stopcodon.
 The CDS has been inserted into pDONR201 via a BP Clonase(TM)
 reaction. Additional sequence has been added in front of the start
 codon: attc...AAAAA GCA GGC TCC ACC (ATG).
 The last codon is followed by the 3' att site: GACCCAGCTTTCTT. att
 The clone is validated by full sequence check.
 Compared to the reference sequence BC014098
 We did not find any amino acid exchanges.
 Clone distribution: http://www.rzpd.de/products/orfclones/
 Location/Qualifiers
 1..381
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="RZPD0834B0231D"
 /clone_lib="Human Full ORF Clones Gateway(TM) - RZPD"
 /lab_host="DH5Alpna"
 /note="Vector: pDONR201, Site_1: attP1, Site_2: attP2"
 1..381
 /gene="SNCG"
 1..>381
 /gene="SNCG"
 /codon_start=1
 /protein_id="CAG46589.1"
 /db_xref="GI:49456537"
 /translation="MDVFKGFSIAKEGVGAVERTKQGVTEAEKTKGVMYVGAKT
 KENVQSTVSVAEKTKQKQANAVSVSVTVAKTYBEANINAVTSGVAKEDLRP
 SAPQOEGASKEKEVAEEAQSQGD"
 ORIGIN
 Alignment Scores:
 Pred. No.: 45.6 Length: 381
 Score: 41.00 Matches: 8
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 9 Gaps: 0
 US-09-017-715A-2_COPY_120_127 (1-8) x CR541790 (1-381)
 Qy 1 Giugluaiaglnsercylglyasp 8
 Db 358 GAGAGAGCCCAAGTCGGGAGAC 381
 RESULT 2
 CR541788 384 bp mRNA linear PRI 29-JUN-2004
 LOCUS CR541788
 DEFINITION Homo sapiens full open reading frame cDNA clone RZPD0834F0930D for
 gene SNCG, synuclein, gamma (breast cancer-specific protein 1);
 complete cds, incl. stopcodon.
 ACCESSION CR541788
 VERSION CR541788.1 GI:49456532
 KEYWORDS Full ORF shuttle clone, Gateway(TM), complete cds.
 SOURCE Homo sapiens (human)

ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE
 AUTHORS
 1 (bases 1 to 384)
 Halleck,A., Ebert,L., Moundinya,M., Schick,M., Eisenstein,S.,
 Neubert,P., Kstrang,K., Schatten,R., Shen,B., Henze,S., Mar,W.,
 Korn,B., Zuo,D., Hu,Y. and Labaer,J.
 Cloning of human full open reading frames in Gateway(TM) system
 entry vector (pDONR201)
 TITLE
 JOURNAL
 Unpublished
 2 (bases 1 to 384)
 Halleck,A., Ebert,L., Moundinya,M., Schick,M., Eisenstein,S.,
 Neubert,P., Kstrang,K., Schatten,R., Shen,B., Henze,S., Mar,W.,
 Korn,B., Zuo,D., Hu,Y. and Labaer,J.
 Direct Submission
 Submitted (28-JUN-2004) RZPD Deutsches Ressourcenzentrum fuer
 Genomforschung GmbH, Im Neuenheimer Feld 580, D-69120 Heidelberg,
 Germany
 COMMENT
 RZPD; RZPD0834F0930D, ORFNo 3605
 www.rzpd.de/cgi-bin/products/cl.cgi?cloneid=RZPD0834F0930D RZPDLIB;
 Human Full ORF Clones Gateway(TM) - RZPD (kan-resist.) RZPD LIB No.
 834
 www.rzpd.de/cgi-bin/products/showlib.pl.cgi/response?libNo=834
 www.rzpd.de/products/orfclones/
 Contact: Inge Axtelt
 RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH,
 Heubnerweg 6, D-14059 Berlin, Germany
 Tel: +49 30 32639 100
 Fax: +49 30 32639 111
 www.rzpd.de
 This clone is available from RZPD;
 Contact RZPD (customer.service@rzpd.de) for further information.
 Clone name at Harvard Institute of Proteomics
 (www.hip.harvard.edu): FLH131021.01X
 This CDS clone is part of a collection of human full ORF clones
 jointly established and verified by the Harvard Institute of
 Proteomics (HIP) and RZPD.
 This CDS has been inserted incl. stopcodon.
 The CDS has been inserted into pDONR201 via a BP Clonase(TM)
 reaction. Additional sequence has been added in front of the start
 codon: attc...AAAAA GCA GGC TCC ACC (ATG).
 The stopcodon is followed by the 3' att site: GACCCAGCTTTCTT. att
 The clone is validated by full sequence check.
 Compared to the reference sequence BC014098
 We did not find any amino acid exchanges.
 Clone distribution: http://www.rzpd.de/products/orfclones/
 Location/Qualifiers
 1..384
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="RZPD0834F0930D"
 /clone_lib="Human Full ORF Clones Gateway(TM) - RZPD"
 /lab_host="DH5Alpna"
 /note="Vector: pDONR201, Site_1: attP1, Site_2: attP2"
 1..384
 /gene="SNCG"
 1..384
 /gene="SNCG"
 /codon_start=1
 /protein_id="CAG46587.1"
 /db_xref="GI:49456533"
 /translation="MDVFKGFSIAKEGVGAVERTKQGVTEAEKTKGVMYVGAKT
 KENVQSTVSVAEKTKQKQANAVSVSVTVAKTYBEANINAVTSGVAKEDLRP
 SAPQOEGASKEKEVAEEAQSQGD"
 ORIGIN
 Alignment Scores:
 Pred. No.: 45.9 Length: 384
 Score: 41.00 Matches: 8
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0

DB: 9 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x CR5411788 (1-384)

QY 1 GluGlulAaInserGlyIyasp 8
Db 358 GAGGAGGCCAGAGTGGGGAGAC 381

RESULT 3
AF411524 488 bp mRNA linear PRI 20-SEP-2001

LOCUS AF411524
DEFINITION Homo sapiens synuclein gamma mRNA, complete cds.
ACCESSION AF411524
VERSION AF411524.1 GI:15705904

KEYWORDS
SOURCE Homo sapiens (human)

ORGANISM Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 488)
AUTHORS Han,C., Zhang,B., Peng,X., Yuan,J. and Qiang,B.
TITLE Direct Submission
JOURNAL Submitted (19-AUG-2001) Department of Biochemistry, Institute of
Basic Medical Science, Chinese Academy of Medical Sciences, 5 Dong
Dan San Tiao, Beijing 100005, P.R. China

FEATURES
source Location/Qualifiers
1..488
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
12..395
/note="SNGC: breast cancer-specific protein 1"

CDS
/codon_start=1
/product="synuclein gamma"
/protein_id="AA105870.1"
/db_xref="GI:15705905"
/translation="MDVFKGFSIAKEGVDAVEKTKGVTEAEKTKGVMYVGAKT
KENVOSTVTSVAKTEKQANAVSAVSVNTATKVEBAENIAVTSSVRRDELARP
SAQQQBEBAKEKEVNAEENQSGD"

ORIGIN

Alignment Scores:

Pred. No.: 56.4 Length: 488
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x AF411524 (1-488)

QY 1 GluGlulAaInserGlyIyasp 8
Db 369 GAGGAGGCCAGAGTGGGGAGAC 392

RESULT 4
AR412236 550 bp DNA linear PAT 18-DEC-2003

LOCUS AR412236
DEFINITION Sequence 1 from patent US 6639052.
ACCESSION AR412236
VERSION AR412236.1 GI:40167022

KEYWORDS
SOURCE Unknown.

ORGANISM Unknown.
Unclassified.
1 (bases 1 to 550)

REFERENCE 1 (bases 1 to 550)
AUTHORS Moore,P.A.
TITLE Human ADA2 polypeptides
JOURNAL Patent: US 6639052-A 1 28-OCT-2003;
FEATURES Location/Qualifiers
1..550
/organism="unknown"
/mol_type="genomic DNA"

ORIGIN

Alignment Scores:

Pred. No.: 62.5 Length: 550
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x AR412236 (1-550)

QY 1 GluGlulAaInserGlyIyasp 8
Db 369 GAGGAGGCCAGAGTGGGGAGAC 392

RESULT 5
AX331171 550 bp DNA linear PAT 09-JAN-2002

LOCUS AX331171
DEFINITION Sequence 1680 from Patent WO0194629.
ACCESSION AX331171
VERSION AX331171.1 GI:18121805

KEYWORDS
SOURCE Homo sapiens (human)

ORGANISM Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1
AUTHORS Young,P.E., Augustus,M., Carter,K.C., Ebner,R., Endress,G.,
Horrigan,S., Soppet,D.R. and Weaver,Z.
TITLE Cancer gene determination and therapeutic screening using signature
gene sets

JOURNAL Patent: WO 0194629-A 1680 13-DEC-2001;
FEATURES Avalon Pharmaceuticals (US)
source Location/Qualifiers
1..550
/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"

ORIGIN

Alignment Scores:

Pred. No.: 62.5 Length: 550
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x AX331171 (1-550)

QY 1 GluGlulAaInserGlyIyasp 8
Db 369 GAGGAGGCCAGAGTGGGGAGAC 392

RESULT 6
BD022727 550 bp DNA linear PAT 27-AUG-2002

LOCUS BD022727
DEFINITION Mammary cancer-specific gene 1.
ACCESSION BD022727
VERSION BD022727.1 GI:22563950

KEYWORDS
SOURCE Homo sapiens (human)

ORGANISM Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 550)
AUTHORS Ji,H. and Rosen,C.A.
TITLE Mammary cancer-specific gene 1
JOURNAL Patent: JP 2001509664-A 1 24-JUL-2001;
COMMENT HUMAN GENOME SCIENCES INC
PN JP 2001509664-A/1
PD 24-JUL-2001

US-09-017-715A-2_COPY_120_127 (1-8) x BD022727 (1-550)

QY 1 GIUGIUAAGIAGIAGIYGLYASP 8

Db 369 GAGGAGGCCCGAGAGTGGGGGAGAC 392

RESULT 7
LOCUS AF010126 550 bp mRNA linear PRI 26-JUL-1997
DEFINITION Homo sapiens breast cancer-specific protein 1 (BCSG1) mRNA,
Homo cds.
ACCESSION AF010126
VERSION AF010126.1 GI:2281473
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 550)
AUTHORS Ji,H., Liu,Y.E., Jia,T., Wang,M., Liu,J., Xiao,G., Joseph,B.K.,
Rosen,C. and Shi,Y.E.
TITLE Identification of a breast cancer-specific gene, BCSG1, by direct
differential cDNA sequencing
JOURNAL Cancer Res. 57 (4), 759-764 (1997) *File 154*
MEDLINE 97178957
REFERENCE 9044857
PUBMED
JOURNAL 2 (bases 1 to 550)
AUTHORS Ji,H., Liu,Y.E., Jia,T., Wang,M., Liu,J., Xiao,G., Joseph,B.K.,
Rosen,C. and Shi,Y.E.
TITLE Direct Submission
JOURNAL Submitted (23-JUN-1997) Ped. Res., Long Island Jewish Medical
Center, 270-05 76th Ave., New Hyde Park, NY 11040, USA
FEATURES
source Location/Qualifiers
1..550
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/tissue_type="breast cancer"
/note="cDNA highly abundant in a breast cancer library but
not in normal library"
1..550
/gene="BCSG1"
12..395
/gene="BCSG1"
/note="breast cancer-specific protein 1, synonym:like;
AD amyloid-like"
/codon_start=1

ORIGIN	<pre> /product="BCG1_protein" /protein_id="AAB64109.1" /db_xref="GI:2281474" /translation="MDVFKFSGIARKGVGAVEKTKOGLTEAAKTEKGMVYGAKTK KENVQSVTSVAEKTKEQANAAVSKAVSSVNTATKIVEAENIATVSGVRRKDLRP SAPOQEGEASKEKEVEAEASQSD" </pre>			
Alignment Scores:				
Pred. No.:	62.5	Length:	550	
Score:	41.00	Matches:	8	
Percent Similarity:	100.00%	Conservative:	0	
Best Local Similarity:	100.00%	Mismatches:	0	
Query Match:	100.00%	Indels:	0	
DB:	9	Gaps:	0	
US-09-017-715A-2_COPY_120_127 (1-8) x AF010126 (1-550)				
CY	1	GLGULNALagInserGIyGIyASP	8	
DB	369	GAGGAGGCCCCAGAGTGGGGGAGAC	392	
RESULT 8				
LOCUS	COJ20882	704 bp	DNA	linear PAT 03-FEB-2004
DEFINITION	Sequence 6816 from Patent WO02068579.			
ACCESSION	COJ20882			
VERSION	COJ20882.1	GI:4281739		
KEYWORDS				
SOURCE	Homo sapiens (human)			
ORGANISM	Homo sapiens			
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.			
TITLE	Venter,C.J., Adams,M.C., Li,P.W. and Myers,E.W.			
	Kits, such as nucleic acid arrays, comprising a majority of			
	humansxons or transcripts, for detecting expression and other uses			
	thereof			
JOURNAL	Patent: WO 02068579-A, 6816 06-SEP-2002;			
	PE Corporation (NY) (US)			
FEATURES	Location/Qualifiers			
source	1..704			
	/organism="Homo sapiens"			
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	/db_xref="taxon:9606"			
ORIGIN	<pre> Alignment Scores: Pred. No.: 77.2 Length: 704 Score: 41.00 Matches: 8 Percent Similarity: 100.00% Conservative: 0 Best Local Similarity: 100.00% Mismatches: 0 Query Match: 100.00% Indels: 0 DB: 6 Gaps: 0 </pre>			
US-09-017-715A-2_COPY_120_127 (1-8) x COJ20882 (1-704)				
CY	1	GLGULNALagInserGIyGIyASP	8	
DB	405	GAGGAGGCCCCAGAGTGGGGGAGAC	428	
RESULT 9				
LOCUS	E36333	720 bp	DNA	linear PAT 18-JUN-2001
DEFINITION	Analytical matter based on synuclein and novel synuclein protein.			
ACCESSION	E36333			
VERSION	E36333.1	GI:13022626		
KEYWORDS	JP 1999239488-A/1.			
SOURCE	Homo sapiens (human)			
ORGANISM	Homo sapiens			
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
	Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.			
REFERENCE	1 (bases 1 to 720)			

AUTHORS Andrew, S.M., Valdimira, R.B. and Arun, M.D.
TITLE Analytical matter based on synuclein and novel synuclein protein
JOURNAL Patent: JP 1999239488-A 1 07-SEP-1999;
COMMENT THE UNIVERSITY COURT OF THE UNIVERSITY OF ST ANDREWS, NUTROBA LTD
OS Homo sapiens (human)
PN JP 1999239488-A/1
PD 07-SEP-1999
PR 21-SEP-1998 JP 1998306283
PI 19-SEP-1997 GB 9719879.0
PT ANDREW SMITH MAKARION, VALDIMIRA RIVOICHI BUCHIMAN, PI ARUN
MLTWARD DAVIS
PC C12N15/09, A01K67/027, C12Q1/68, G01N33/53, C12N15/00 CC
FI Key Location/Qualifiers
FT CDS (49) . (432).
FEATURES Location/Qualifiers
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ORIGIN /organism="Homo sapiens"
Alignment Scores: /mol_type="genomic DNA"
Pred. No.: /db_xref="taxon:9606"
Score: 78.7 Length: 720
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Best Local Similarity: 100.00% Conservative: 0
Query Match: 100.00% Mismatches: 0
DB: Indels: 0
US-09-017-715A-2_COPY_120_127 (1-8) x E36333 (1-720)
QY 1 GluGlualaglnSerglygIyasp 8
Db 406 GAGGAGGCCAGAGTGGGGAGAC 429
RESULT 10
AX004527 AX004527 720 bp DNA linear PAT 24-AUG-2000
LOCUS Sequence 1 from Patent EP0908727.
DEFINITION AX004527
ACCESSION AX004527.1 GI:9927977
VERSION
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
REFERENCE 1 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
AUTHORS Synuclein-based assay and synuclein protein
TITLE Patent: EP 0908727-A 1 14-APR-1999;
JOURNAL NEUROPA LIMITED (GB); UNIV COURT OF THE UNIVERSITY O (GB)
FEATURES Location/Qualifiers
Source 1..720
ORIGIN /organism="Homo sapiens"
Alignment Scores: /mol_type="unassigned DNA"
Pred. No.: /db_xref="taxon:9606"
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Query Match: 100.00% Mismatches: 0
DB: Indels: 0
US-09-017-715A-2_COPY_120_127 (1-8) x AF017256 (1-720)
QY 1 GluGlualaglnSerglygIyasp 8
Db 406 GAGGAGGCCAGAGTGGGGAGAC 429
RESULT 12
LOCUS BVL177827 738 bp DNA linear STS 10-JUN-2004
DEFINITION sqm97020 Human DNA (Sequencm) Homo sapiens STS genomic, sequence
ACCESSION BVL177827
VERSION BVL177827.1 GI:48014020

US-09-017-715A-2_COPY_120_127 (1-8) x AX004527 (1-720)
QY 1 GluGlualaglnSerglygIyasp 8
Db 406 GAGGAGGCCAGAGTGGGGAGAC 429
RESULT 11
AF017256 AF017256 720 bp mRNA linear PRI 23-SEP-1998
LOCUS Homo sapiens peryn mRNA, complete cds.
DEFINITION AF017256
ACCESSION AF017256.1 GI:3642774
VERSION
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
REFERENCE 1 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
AUTHORS Minkina, N.N., Alimova-Kost, M.V., Paterson, J.W., Delaney, L.,
TITLE Cohen, B.B., Imreh, S., Gnuchey, N.V., Davies, A.M., and Buchman, V.L.
JOURNAL Hum. Mol. Genet. 7 (9), 1417-1424 (1998)
MEDLINE 98367030
PUBMED 9700196
REFERENCE 2 (bases 1 to 720)
AUTHORS Buchman, V.L.
TITLE Direct Submision
JOURNAL Submitted (04-AUG-1997) School of Biomedical Sciences, Univ. of St.
ANDREWS Bute Medical Buildings, St. Andrews, Fife KY16 9TS,
SCOTLAND
FEATURES Location/Qualifiers
Source 1..720
ORIGIN /organism="Homo sapiens"
Alignment Scores: /mol_type="mRNA"
Pred. No.: /db_xref="taxon:9606"
Score: 49.432 Length: 720
Percent Similarity: 100.00% Matches: 8
Best Local Similarity: 100.00% Conservative: 0
Query Match: 100.00% Mismatches: 0
DB: Indels: 0
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QY 1 GluGlualaglnSerglygIyasp 8
Db 406 GAGGAGGCCAGAGTGGGGAGAC 429
RESULT 12
LOCUS BVL177827 738 bp DNA linear STS 10-JUN-2004
DEFINITION sqm97020 Human DNA (Sequencm) Homo sapiens STS genomic, sequence
ACCESSION BVL177827
VERSION BVL177827.1 GI:48014020

Thu May 5 15:10:43 2005

us-09-017-715a-2_copy_120_127.rge

Page 9

Db 223025 GAGAGCACAGAGTGGAGGGGAC 223048

Search completed: May 4, 2005, 11:54:02
Job time : 489.669 secs

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OM protein - nucleic search, using frame_plus_p2n model

Run on: May 4, 2005, 09:07:55 ; Search time 7212.58 Seconds
(without alignments)
853.206 Million cell updates/sec

Title: US-09-017-715A-2

Perfect score: 610
Sequence: 1 MDVFKGKFSIAKKGVGAVE.....EGEASKEKEVAEHAQSGSD 127

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Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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-UNITS=bits -START=1 -END=1 -MATRIX=blonum62 -TRANS=human40.cdi -LIST=45
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-OUTFMT=ptc -NOR=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=200000000
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-DEV TIMEOUT=120 -MAIN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

- GenEmbl:*
- 1: gb_ba:*
- 2: gb_hvg:*
- 3: gb_in:*
- 4: gb_cm:*
- 5: gb_ov:*
- 6: gb_pat:*
- 7: gb_ph:*
- 8: gb_pl:*
- 9: gb_pr:*
- 10: gb_ro:*
- 11: gb_sts:*
- 12: gb_sy:*
- 13: gb_un:*
- 14: gb_vi:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	610	100.0	550	AR412236	AR412236 Sequence
2	610	100.0	550	AX331171	AX331171 Sequence
3	610	100.0	550	BD022727	BD022727 Mammary c
4	610	100.0	550	AF010126	AF010126 Homo sapi

5	602	98.7	381	9	CR541790	CR541790 Homo sapi
6	602	98.7	384	9	CR541788	CR541788 Homo sapi
7	602	98.7	704	6	CQ720882	CQ720882 Sequence
8	602	98.7	758	9	BC014098	BC014098 Homo sapi
9	595	97.5	488	9	AF411524	AF411524 Homo sapi
10	595	97.5	720	6	E36333	E36333 Analytical
11	595	97.5	720	6	AX004527	AX004527 Sequence
12	595	97.5	720	9	AF017256	AF017256 Homo sapi
13	532.5	87.3	738	11	BV177827	BV177827 sqm97020
14	523	85.7	677	4	AF219257	AF219257 Bos tauru
15	497	81.5	727	6	E36334	E36334 Analytical
16	497	81.5	727	6	AX004529	AX004529 Sequence
17	497	81.5	727	6	AF017255	AF017255 Mus muscu
18	497	81.5	748	10	BC028508	BC028508 Mus muscu
19	476	78.0	488	10	AY518351	AY518351 Rattus no
20	470	77.0	732	10	RNSDSYNGE	X86789 R. norvegicu
21	438.5	71.9	873	5	AF253513	AF253513 Gallus ga
22	412	67.5	1120	5	CR762140	CR762140 Xenopus t
23	399.5	65.5	1046	5	BC054269	BC054269 Xenopus l
24	388.5	63.7	1040	5	AY055119	AY055119 Xenopus l
25	388.5	63.7	1180	5	BC072217	BC072217 Xenopus l
26	316	51.8	695	10	S73007	S73007 synuclein S
27	316	51.8	1017	10	AY550005	AY550005 Rattus no
28	316	51.8	1017	10	AY550006	AY550006 Rattus no
29	316	51.8	1018	10	AF007758	AF007758 Rattus no
30	316	51.8	1047	10	BC046764	BC046764 Mus muscu
31	316	51.8	1124	10	AF033261	AF033261 Mus muscu
32	316	51.8	1181	10	AF179273	S73008 synuclein S
33	315	51.6	695	10	S73008	L33860 Serinus can
34	314.5	51.6	1234	5	SEISYNELFI	AF253512 Rattus no
35	311.5	51.1	1182	5	AF253512	AF253512 Gallus ga
36	311.5	51.1	1190	5	EX936116	EX936116 Gallus ga
37	311	51.0	1233	5	BC054200	BC054200 Xenopus l
38	310	50.8	549	10	AF044672	AF044672 Mus muscu
39	308	50.5	549	10	RAT14KXAP	D17764 Rattus sp.
40	307.5	50.4	420	9	CR541653	CR541653 Homo sapi
41	307.5	50.4	423	6	AX662310	AX662310 Sequence
42	307.5	50.4	423	9	CR457058	CR457058 Homo sapi
43	307.5	50.4	423	12	BT007765	BT007765 Synthetic
44	307.5	50.4	425	9	AY049786	AY049786 Homo sapi
45	307.5	50.4	546	6	CQ728889	CQ728889 Sequence

ALIGNMENTS

RESULT 1
AR412236
LOCUS AR412236 550 bp DNA
DEFINITION Sequence 1 from patent US 6639052.
ACCESSION AR412236
VERSION AR412236.1 GI:40167022
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 550)
AUTHORS Moore, P.A.
TITLE Human ADA2 polypeptides
JOURNAL Patent: US 6639052-A 1 28-OCT-2003;
FEATURES
source 1..550
/organism="unknown"
/mol_type="genomic DNA"

ORIGIN

Alignment Scores:
Pred. No.: 7.25e-44
Score: 610.00
Percent Similarity: 100.00%
Best Local Similarity: 100.00%
Query Match: 100.00%
DB: 6
Length: 550
Matches: 127
Conservative: 0
Mismatch: 0
Indels: 0
Gaps: 0

US-09-017-715A-2 (1-127) x AR412236 (1-550)

QY 1 MetaspvalPheLysGlyPheSerIleAlaLysGlyValValGluValGlu 20
DB 12 ATGAGATGTTTCAAGAGGCGCTTCTCCATCGCCAAAGGCGGTGGGTGGGAGAA 71
QY 21 LysThrLysGlnGlyValThrGluAlaGluLysThrLysGluGluValMetCysVal 40
DB 72 AAGACCAACAGAGGGGTGACGGAAGCAGCTGAGAAACCAAGAGGGGTGCATGATGTG 131
QY 41 GtAlaLysThrLysGluAsnValValGlnSerValThrSerValAlaGluLysThrLys 60
DB 132 GGAGCCAAAGCAGAGGATGTTGTACAGAGCGTACCTCGGGTGGCCAGAGCAAG 191
QY 61 GtGlnAlaAsnAlaValSerLysAlaValSerSerValAsnThrValAlaThrLys 80
DB 192 GAGCAGGCCAAGCGCGTGGAGCAAGGCTGTGTGAGCAGGCTCAACCTGTGGCCACCAAG 251
QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValValArgLysGluAsnLeu 100
DB 252 ACCGTGAGAGAGCGGAGAACATCGCGTCACTCGGGGTGGTGGCAAGAGGACTTG 311
QY 101 ArgProSerAlaProGlnGlnGluGluAlaSerLysGluLysGluGluValAlaGlu 120
DB 312 AGGCCATCTGCCCGCCCAAGAGGCGGTGAGCATCCAAAGAGAGAGAGTGGCAGAG 371
QY 121 GtAlaGlnSerGlyGlyAsp 127
DB 372 GAGGCCAAGAGTGGGAGAGC 392

RESULT 2
AX331171
LOCUS AX331171 550 bp DNA linear PAT 09-JAN-2002
DEFINITION Sequence 1680 from Patent WO0194629.
ACCESSION AX331171
VERSION AX331171.1 GI:18121805
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1
AUTHORS Young, P. E., Augustus, M., Carter, K. C., Ebner, R., Endress, G.,
Hortigan, S., Soppet, D. R. and Weaver, Z.
TITLE Cancer gene determination and therapeutic screening using signature
gene sets
JOURNAL Patent: WO 0194629-A, 1680 13-DEC-2001;
FEATURES Avalon Pharmaceuticals (US)
source 1. 550 Location/Qualifiers
/organism="Homo sapiens"
/mol_type="unassigned DNA"
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ORIGIN

Alignment Scores:
Pred. No.: 7, 25e-44 Length: 550
Score: 610.00 Matches: 127
Percent Similarity: 100.00% Conservative: 0
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DB: 6 Gaps: 0

US-09-017-715A-2 (1-127) x AX331171 (1-550)

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QY 21 LysThrLysGlnGlyValThrGluAlaGluLysThrLysGluGluValMetCysVal 40
DB 72 AAGACCAACAGAGGGGTGACGGAAGCAGCTGAGAAACCAAGAGGGGTGCATGATGTG 131

QY 41 GtAlaLysThrLysGluAsnValValGlnSerValThrSerValAlaGluLysThrLys 60
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QY 61 GtGlnAlaAsnAlaValSerLysAlaValSerSerValAsnThrValAlaThrLys 80
DB 192 GAGCAGGCCAAGCGCGTGGAGCAAGGCTGTGTGAGCAGGCTCAACCTGTGGCCACCAAG 251
QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValValArgLysGluAsnLeu 100
DB 252 ACCGTGAGAGAGCGGAGAACATCGCGTCACTCGGGGTGGTGGCAAGAGGACTTG 311
QY 101 ArgProSerAlaProGlnGlnGluGluAlaSerLysGluLysGluGluValAlaGlu 120
DB 312 AGGCCATCTGCCCGCCCAAGAGGCGGTGAGCATCCAAAGAGAGAGAGTGGCAGAG 371
QY 121 GtAlaGlnSerGlyGlyAsp 127
DB 372 GAGGCCAAGAGTGGGAGAGC 392

RESULT 3
BD022727
LOCUS BD022727 550 bp DNA linear PAT 27-AUG-2002
DEFINITION Mammary cancer-specific gene 1.
ACCESSION BD022727
VERSION BD022727.1 GI:22563950
KEYWORDS JP 2001509664-A/1.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1
AUTHORS Ji, H. and Rosen, C. A.
TITLE Mammary cancer-specific gene 1
JOURNAL Patent: JP 2001509664-A, 1 24-JUL-2001;
COMMENT HUMAN GENOME SCIENCES INC
PN JP 2001509664-A/1
PD 24-JUL-2001
PF 03-FEB-1998 JP 1998515053
PR 03-FEB-1997 US 60/037080
PI HONGJUN JI, CRAIG A ROSEN

PC C12N15/09, C07K14/47, C07K16/18, C12N1/15, C12N1/19, C12N1/21, C12N5/ PC
10,
PC C12P21/02, C12Q1/68, G01N33/574, C12N15/00, C12N5/00 CC
Strandedness: Double;
CC Topology: Both; Location/Qualifiers
FH Key 12. 392.
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source 1. 550
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"

ORIGIN

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Pred. No.: 7, 25e-44 Length: 550
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Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-09-017-715A-2 (1-127) x BD022727 (1-550)

QY 1 MetaspvalPheLysGlyPheSerIleAlaLysGlyValValGluValGlu 20
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QY 21 LysThrLysGlnGlyValThrGluAlaGluLysThrLysGluGluValMetCysVal 40
DB 72 AAGACCAACAGAGGGGTGACGGAAGCAGCTGAGAAACCAAGAGGGGTGCATGATGTG 131

OY		41	GlyAlaValThrLySGluAsnValGlnSerValThrSerValAlaGluLysThrlys 	60
Dd		132	GGAGCGAAGAACCACGAAGAATGTATTACAAGACCGTCACTGTGCCGAGAAAGCAAAG 	191
OY		61	GluGlnAlaAsnAlaValSerIySerylaValSerSeServalasnThrValAlathrlrys 	80
Dd		192	GAGCAGGCACACCCGTGTAGCACAGGCTGTGGTGACACGTCATAACTGTGCCAACCAAG 	251
OY		81	ThrValGlUGlUalAGluAsnllealaValThrSerGIylvalValArXyluglInspleu 	100
Dd		252	ACCGTGGAGGAGGCGGAGAACACTCCCGGTCACTCCGGGGTGGTCCGACAGAGCATTTG 	311
OY		101	ArgProserAlaproGlngInglngluLygLuJaserylsglutryglugluValalalu 	120
Dd		312	AGGCCATCTGCCCCCCAACAGAGGGGTAGGATCCAAGAAAAGAGAAAGTAGGACAGAG 	371
OY		121	GluAlaglnSerGIylGLYASP 	127
Dd		372	GAGGCCACAGATGTGGGGAGAC 	392
RESULT 4				
LOCUS	AF010126	550 bp mRNA linear PRI 26-JUL-1997		
DEFINITION	Homo sapiens breast cancer-specific protein 1 (BCSG1) mRNA,			
complete cds.				
VERSION	AF010126			
KEYWORDS	AF010126.1 GI:2281473	Feb 15th		
SOURCE	Homo sapiens (human)			
ORGANISM	Homo sapiens			
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;				
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.				
Ji.H., Liu,Y.E.; Jia,T., Wang,M., Liu,J., Xiao,G., Joseph,B.K.,				
Rosen,C. and Shi.Y.E.				
Identification of a breast cancer-specific gene, BCSG1, by direct				
differential CDNA sequencing				
Cancer Res. 57 (4), 759-764 (1997)				
TITLE	Journal MEDLINE			
PUBMED	97178957			
REFERENCE	2 (bases 1 to 550)			
AUTHORS	JI,H., LIU,Y.E., JIA,T., WANG,M., LIU,J., XIAO,G., JOSEPH,B.K., ROSEN,C. AND SHI,Y.E.			
TITLE	Direct Submission			
JOURNAL	Submitted (23-JUN-1997) Ped. Res., Long Island Jewish Medical Center, 270-05 /6th Ave., New Hyde Park, NY 11040, USA			
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/feature_type="breast cancer"				
/note="CDNA highly abundant in a breast cancer library but not in normal library"				
1..550				
/gene="BCSG1"				
12..395				
/gene="BCSG1"				
/note="breast cancer-specific protein 1; synuclein-like; AD amyloid-like"				
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/product="BCSG1 protein"				
/protein_id="BAB64109.1"				
/db_xref="gi:2281474"				
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ORIGIN				
Alignment Scores:	7.25e+44	Length:	550	
Pred. No.:	610..00	Matches:	127	
Score:				

	Percent Similarity:	100.00%	Conservative:	0
	Best Local Similarity:	100.00%	Mismatches:	0
	Query Match:	100.00%	Indels:	0
DB:		9	Gaps:	0
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OY		1 MetaspvAlPheIySvlglyPhSeSerlleaIalyslvsglyValalGlvalAvalGlu	20	
Dd		12 ATGATGTTTTCAAGAAAGGGCTTCCATGCCCAAGAAGGGCGTGCGGTGGAA	71	
OY		21 LvsThrlysgInglYvalThrGUaAlaAgIulvyrThrlysgInglYvalMeTyrcVal	40	
Dd		72 AAGACCAGACGGGGGGTGA CGAACGCAGCTGAGAAACCAAGAGGGGGTATGATGTG	131	
OY		41 GlYlaIayThrlysgIuaSnVlalGlInserValThrservalalagIulvSThrlys	60	
Dd		132 GGAGCCAAAGACCAAGAGAAATTGTGTACAAGCGCTGACTAGTGGCCGAAGCCAA	191	
OY		61 GlUGlnAlaaenAlavalserlyslavalvalserseValaanThrvallalThrlys	80	
Dd		192 GAGCAGGCCCAACGCCGTGACCAAGCGTGTGTGAGCAGCTCAACACTGTGGCCACCA	251	
OY		81 ThrValGluGuIualaglUasnlIalavalThrseryValvalaclyvgIuaspleu	100	
Dd		252 ACCGTGGAGGAGGGCGGAACATGCGCATCCCTCCGGGGTGGTCGCAAGAGGACTTG	311	
OY		101 ArgProSerAlaPCoGInglnglUGlYlalaSerlysgIulvsgIuValalaglu	120	
Dd		312 AGCGCATCTGCCCCCCCACAGAGAGGTGTGAGCATCCAAGAAAGAGGAGTGGCAG	371	
OY		121 GlUaIagInserygIylap 127		
Dd		372 GAGGCCAGAGATGGGGGAGAC 392		
 RESULT 5				
CRS41790				
LOCUS				
DEFINITION		Homo sapiens full open reading frame cDNA clone RZPD0834B0231D for gene SNGC, myonuclein, gamma (breast cancer-specific protein 1); complete cds, without stopcodon.		
ACCESSION		CRS41790		
VERSION		CRS41790.1		
KEYWORDS		Full ORF shuttle clone, Gateway(TM), complete cds.		
SOURCE		Homo sapiens (human)		
ORGANISM		Homo sapiens		
REFERENCE		Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.		
AUTHORS		1 (bases 1 to 381) Halleck,A., Ebert,L., Mkoundinya,M., Schick,M., Bisenstein,S., Neubert,P., Ketrang,K., Schatten,R., Shen,B., Henze,S., Mar,W., Korn.B., Zuo,D., Hu.Y. and Labaer,J.		
TITLE		Cloning of human full open reading frames in Gateway(TM) system entry vector (pDONR201)		
JOURNAL		Unpublished		
REFERENCE		2 (bases 1 to 381) Halleck,A., Ebert,L., Mkoundinya,M., Schick,M., Bisenstein,S., Neubert,P., Ketrang,K., Schatten,R., Shen,B., Henze,S., Mar,W., Korn.B., Zuo,D., Hu.Y. and Labaer,J.		
AUTHORS		Direct Submission Submitted (28-JUN-2004) RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH, Im Neuenheimer Feld 580, D-69120 Heidelberg, Germany		
TITLE		RZPD: RZPD0834B0231D, ORFno 3631		
JOURNAL		www.rzpd.de/cgi-bin/products/cl.cgi?cloneID=RZPD0834B0231D RZPDLIB: Human Full ORF Clones Gateway(TM) - RZPD (kan-resistc.) RZPD LIB NO. 834		
COMMENT		www.rzpd.de/cgi-bin/products/showlib.pl.cgi/?response?libNo=834 RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH, Heubnerweg 6, D-14059 Berlin, Germany Tel.: +49 30 32639 100		

Fax: +49 30 32639 111
 www.rzpd.de
 This clone is available from RZPD:
 Contact RZPD (customer.service@rzpd.de) for further information.
 Clone name at Harvard Institute of Proteomics
 (www.hip.harvard.edu): FLH130940.01L
 This CDS clone is part of a collection of human full ORF clones
 jointly established and verified by the Harvard Institute of
 Proteomics (HIP) and RZPD.
 This CDS has been cloned without stopcodon.
 The CDS has been inserted into pDONR201 via a BP Clonase(TM)
 reaction. Additional sequence has been added in front of the start
 codon: attc. AAAA GCA GGC TCC ACC (ATG).
 The last codon is followed by the 3' att site: GACCCAGCTTCTT. att
 The clone is validated by full sequence check.
 Compared to the reference sequence BC014098
 we did not find any amino acid exchanges.
 Clone distribution: http://www.rzpd.de/products/orfclones/
 Location/Qualifiers

FEATURES

source

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1. 381
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="RZPD0834B0231D"
/clone_1ib="Human Full ORF Clones Gateway(TM) - RZPD"
/lab_host="DH5Alpha"
/notes="Vector: pDONR201, Site_1: attPl, Site_2: attp2"
1..381
/gene="SNCG"
1..381
/gene="SNCG"
/codon_start=1
/protein_id="CAG46589.1"
/db_xref="GI:49456537"
/translation="MDVFRKFSIAKEGVAVGAVETKQGVTEAKTEKGVNYGAKT
KENVQSVTAETKEQANAVSEAVSVNTVATKTYVEAENIATVSGVAREDLRP
SAPOQEGEASKEKEVEAEVAGSGD"

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ORIGIN

Alignment Scores:

Pred. No.: 2,38e-43 Length: 381
 Score: 602.00 Matches: 125
 Percent Similarity: 100.00% Conservative: 2
 Best Local Similarity: 98.43% Mismatches: 0
 Query Match: 98.69% Indels: 0
 DB: 9 Gaps: 0

US-09-017-715A-2 (1-127) x CRS41790 (1-381)

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Oy 1 MetAspValPheIysGlyPheSerIleAlaIysGlyValValAlaGluValGlu 20
Db 1 ATGAGTCTTTCAGAGAGGCTTCTCCTCCTCCAGAGAGGCGTGTGGTGGCGAGAA 60
Oy 21 LysThrLysGlnGlyValThrGluAlaGluValThrLysGlnGlyValMetIyrVal 40
Db 61 AAGACCAACAGAGGGGTGACGAGACGCTGAGAACCAAGAGAGGGGTGATGTG 120
Oy 41 GlyAlaLysThrLysGlnValValGlnSerValThrSerValAlaGluLysThrLys 60
Db 121 GGAGGCCAACAGAGAGATGTTGACAGAGGCTCACTGCGCCAGAGAACCAAG 180
Oy 61 GluGlnAlaAsnAlaValSerLysValValValSerSerValAsnThrValAlaThrLys 80
Db 181 GAGCAGGCCAACCCCTGAGCGAGGCTGTGGTACAGCCTCAACACTGTGGCCACCAAG 240
Oy 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValValArgLysGluAspLeu 100
Db 241 ACCGTGAGAGAGCGGAGAACATCGCGTCACTCCGGGGTGTGCGCAAGAGGACTTG 300
Oy 101 ArgProSerAlaProGlnGlnGlnGluGluAlaSerLysGluLysGluValAlaGlu 120
Db 301 AGGCCATCTGCCCCCAAGAGAGGGGTGAGGATCCAAAGAGAAAGAGAAAGTGCAGAG 360

```

Oy 121 GualagInserGlyIyAsp 127
 Db 361 GAGGCCAGAGTGGGGAGAGAC 381

RESULT 6 384 bp mRNA linear PRI 29-JUN-2004
 LOCUS CRS41788
 DEFINITION Homo sapiens full open reading frame CDNA clone RZPD0834F0930D for
 gene SNCG, synclelin, gamma (breast cancer-specific protein 1);
 complete cds, incl. stopcodon.

ACCESSION CRS41788
 VERSION CRS41788.1 GI:49456532
 KEYWORDS Full ORF shuttle clone, Gateway(TM), complete cds.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 384)
 Halleleck,A., Ebert,L., Mroundinya,M., Schick,M., Eisenstein,S.,
 Neubert,P., Ketrang,K., Schatten,R., Shen,B., Henze,S., Mar,W.,
 Korn,B., Zuo,D., Hu,Y. and Labber,J.
 Cloning of human full open reading frames in Gateway(TM) system
 entry vector (pDONR201)
 Unpublished
 2 (bases 1 to 384)
 Halleleck,A., Ebert,L., Mroundinya,M., Schick,M., Eisenstein,S.,
 Neubert,P., Ketrang,K., Schatten,R., Shen,B., Henze,S., Mar,W.,
 Korn,B., Zuo,D., Hu,Y. and Labber,J.
 Direct Submission
 Submitted (28-JUN-2004) RZPD Deutsches Ressourcenzentrum fuer
 Genomforschung GmbH, Im Neuenheimer Feld 580, D-69120 Heidelberg,
 Germany

TITLE JOURNAL
 REFERENCE
 AUTHORS

TITLE JOURNAL
 REFERENCE
 AUTHORS

COMMENT RZPD: RZPD0834F0930D, ORFNo 3605
 www.rzpd.de/cgi-bin/products/cl.cgi?cloneID=RZPD0834F0930D RZPD LIB:
 Human Full ORF Clones Gateway(TM) - RZPD (ken-resistc.) RZPD LIB No.
 834
 www.rzpd.de/cgi-bin/products/showlib.pl.cgi?response?libNo=834
 www.rzpd.de/products/orfclones/
 Contact: Inge Axlart
 RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH,
 Heubnerweg 6, D-14059 Berlin, Germany
 Tel: +49 30 32639 100
 Fax: +49 30 32639 111
 www.rzpd.de

This clone is available from RZPD;
 Contact RZPD (customer.service@rzpd.de) for further information.
 Clone name at Harvard Institute of Proteomics
 (www.hip.harvard.edu): FLH130940.01X
 This CDS clone is part of a collection of human full ORF clones
 jointly established and verified by the Harvard Institute of
 Proteomics (HIP) and RZPD.
 This CDS has been cloned incl. stopcodon.
 The CDS has been inserted into pDONR201 via a BP Clonase(TM)
 reaction. Additional sequence has been added in front of the start
 codon: attc. AAAA GCA GGC TCC ACC (ATG).
 The stopcodon is followed by the 3' att site: GACCCAGCTTCTT. att
 The clone is validated by full sequence check.
 Compared to the reference sequence BC014098
 we did not find any amino acid exchanges.
 Clone distribution: http://www.rzpd.de/products/orfclones/
 Location/Qualifiers

FEATURES

source

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1. 384
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="RZPD0834F0930D"
/clone_1ib="Human Full ORF Clones Gateway(TM) - RZPD"
/lab_host="DH5Alpha"
/notes="Vector: pDONR201, Site_1: attPl, Site_2: attp2"
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/codon_start=1
/protein_id="CA646587.1"
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KENVQSVTSVAEKTKEQANAVSEAVSVNTATKTVEEANIATVSGVVRKEDLRP
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ORIGIN

Alignment Scores: 2.4e-43 Length: 384
Score: 602.00 Matches: 125
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 98.43% Mismatches: 0
Query Match: 98.69% Indels: 0
DB: 9 Gaps: 0

US-09-017-715a-2 (1-127) x CR541788 (1-384)

Qy 1 MetAspValPheIlyblysgIlyPheSerIleAlaIlyblysgIlyValIglValIglu 20
Db 1 ATGATGTCTTCAAGAGGCTTCTCCATGCCCAAGAGGCGTGTGGTCCGATGGA 60
Qy 21 LysThrIybsgIlyValIThrGluAlaIgluIlyblysgIlyValMetTyVal 40
Db 61 AAGACCAAGCAGGGGTGACGGAAGCAGCTGAGAAGCAAGAGGGGTGATGATGTG 120
Qy 41 G1yAlaIlyblysgIlyblyblyblyblyblyblyblyblyblyblyblyblybly 60
Db 121 GAGGCCAAGCAGGAGAAATGTGTACAGAGCTGACCTCAGTGGCCGAGAACCAAG 180
Qy 61 GluGlnAlaAsnAlaValSerIyAlaValIlySerValAsnThrValAlaThrIys 80
Db 181 GAGCAGGCCAAGCGCTGTGAGCAGGCTGTGACAGCTCACTGTGGCCACCAAG 240
Qy 81 ThrValIgluIlyblyblyblyblyblyblyblyblyblyblyblyblyblybly 100
Db 241 ACCGTGAGAGAGGGGAGAACATCGCGTCACTCCGGGGTGTGGCCAGAGGACTTG 300
Qy 101 ArgProSerAlaProGInGInGInGInGInGInGInGInGInGInGInGInGIn 120
Db 301 AGGCATCTGCCCCCAAGAGAGGTGAGGATCCCAAGAGAAAGAGAGTGGCAGAG 360
Qy 121 GluAlaGInSerIyGlyIyAsp 127
Db 361 GAGGCCAGAGTGGGAGAGAC 381

RESULT 7
LOCUS CQ720882 704 bp DNA linear PAT 03-FEB-2004
DEFINITION Sequence 6816 from Patent WO02068579.
ACCESSION CQ720882
VERSION CQ720882.1 GI:42281739
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE
1 Venter,C.J., Adams,M.C., Li,P.W. and Myers,E.W.
Kits, such as nucleic acid arrays, comprising a majority of
humans or transcripts, for detecting expression and other uses
thereof
Patent: WO 02068579-A 6816 06-SEP-2002;

JOURNAL
PE Corporation (NY) (US)
Location/Qualifiers

FEATURES
source
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/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"

ORIGIN
Alignment Scores: 4.59e-43 Length: 704
Pred. No.:

Score: 602.00 Matches: 125
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 98.43% Mismatches: 0
Query Match: 98.69% Indels: 0
DB: 6 Gaps: 0

US-09-017-715a-2 (1-127) x CQ720882 (1-704)

Qy 1 MetAspValPheIlyblysgIlyPheSerIleAlaIlyblysgIlyValIglValIglu 20
Db 48 ATGATGTCTTCAAGAGGCTTCTCCATGCCCAAGAGGCGTGTGGTCCGATGGA 107
Qy 21 LysThrIybsgIlyValIThrGluAlaIgluIlyblysgIlyValMetTyVal 40
Db 108 AAGACCAAGCAGGGGTGACGGAAGCAGCTGAGAAGCAAGAGGGGTGATGATGTG 167
Qy 41 G1yAlaIlyblysgIlyblyblyblyblyblyblyblyblyblyblyblyblybly 60
Db 168 GAGGCCAAGCAGGAGAAATGTGTACAGAGCTGACCTCAGTGGCCGAGAACCAAG 227
Qy 61 GluGlnAlaAsnAlaValSerIyAlaValIlySerValAsnThrValAlaThrIys 80
Db 228 GAGCAGGCCAAGCGCTGTGAGCAGGCTGTGACAGCTCACTGTGGCCACCAAG 287
Qy 81 ThrValIgluIlyblyblyblyblyblyblyblyblyblyblyblyblyblybly 100
Db 288 ACCGTGAGAGAGGGGAGAACATCGCGTCACTCCGGGGTGTGGCCAGAGGACTTG 347
Qy 101 ArgProSerAlaProGInGInGInGInGInGInGInGInGInGInGInGInGIn 120
Db 348 AGGCATCTGCCCCCAAGAGAGGTGAGGATCCCAAGAGAAAGAGAGTGGCAGAG 407
Qy 121 GluAlaGInSerIyGlyIyAsp 127
Db 408 GAGGCCAGAGTGGGAGAGAC 428

RESULT 8
LOCUS BC014098 758 bp mRNA linear PRI 29-JUN-2004
DEFINITION Homo sapiens synuclein, gamma (breast cancer-specific protein 1),
mRNA (cDNA clone MGC:20132 IMAGE:4546444), complete cds.
ACCESSION BC014098
VERSION BC014098.2 GI:33878363
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE
AUTHORS Strausberg,R.L., Peltingold,E.A., Grouse,L.H., Derge,J.G.,
Klausner,R.D., Collins,F.S., Wagner,L., Shenmen,C.M., Schuler,G.D.,
Altschul,S.F., Zeeberg,B., Bueltow,K.H., Schaefer,C.F., Bhat,N.K.,
Hopkins,R.F., Jordan,H., Moore,T., Max,S.I., Wang,J., Hsieh,F.,
Diatchenko,L., Marusina,K., Farmer,A.A., Rubin,G.M., Hong,L.,
Stapleton,M., Soares,M.B., Bonaldo,M.F., Casavant,T.L.,
Schetz,T.E., Brownstein,M.U., Utsid,T.B., Toshiyuki,S.,
Carninci,P., Prange,C., Kaha,S.S., Loquelliano,N.A., Peters,G.J.,
Adams,N.R.D., Mullaby,S.J., Bosak,S.A., McEwan,P.J.,
McKernan,K.J., Malek,J.A., Gunaratne,P.H., Richards,S.,
Worley,K.C., Hale,S., Garcia,A.M., Gay,L.J., Huilyk,S.W.,
Villalón,D.K., Muzny,D.M., Sodergren,E.J., Lu,X., Gibbs,R.A.,
Fahy,J., Helton,E., Kettelman,M., Madan,A., Rodriguez,S.,
Sanchez,A., Whiting,M., Madan,A.C., Young,A.C., Shevchenko,Y.,
Boutard,G.G., Blakesley,R.W., Touchman,J.W., Green,E.D.,
Dickson,M.C., Rodriguez,A.C., Grimwood,J., Schmutz,J., Myers,R.M.,
Butterfield,Y.S., Krzywinski,M.I., Skalski,U., Smalins,D.E.,
Schnerch,A., Schein,J.E., Jones,S.J. and Marra,M.A.
Generation and initial analysis of more than 15,000 full-length
human and mouse cDNA sequences
Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
JOURNAL
PUBMED
12477932
2 (bases 1 to 758)
REFERENCE
AUTHORS Strausberg,R.

TITLE Direct Submission
JOURNAL Submitted (10-SEP-2001) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
USA

REMARK NIH-MGC Project URL: <http://mgc.nci.nih.gov>
On Aug 19, 2003 this sequence version replaced gi:15559464.
COMMENT Contact: MGC help desk
Email: cgabs-remail.nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: National Institutes of Health Intramural
Sequencing Center (NISC),
Gaithersburg, Maryland;
Web site: <http://www.nisc.nih.gov/>
Contact: nisc_mgc@nigrl.nih.gov
Akhter, N., Ayale, K., Beckstrom-Sternberg, S.M., Benjamin, B.,
Blakesley, R.W., Bouffard, G., Breen, K., Brinkley, C., Brooks, S.,
Dierich, N.L., Granite, S., Guan, X., Gupta, J., Haghighi, P.,
Hansen, N., Ho, S.-L., Karlins, E., Kwong, P., Laric, P., Legaspi, R.,
Maduro, Q.L., Mastello, C., Maskeri, B., Mastrian, S.D., McCloskey, J.C.,
McDowell, J., Pearson, R., Stantipop, S., Thomas, P.D., Touchman, J.W.,
Tsugeon, C., Vogt, J.L., Walker, M.A., Wetherby, K.D., Wiggins, L.,
Young, A., Zhang, L.-H. and Green, E.D.

FEATURES
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/clone_id="NIH MGC_15"
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ORIGIN

Alignment Scores:

Pred. No.: 4,97e-43 Length: 758
Score: 602.00 Matches: 125
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 98.43% Mismatches: 0
Query Match: 98.69% Indels: 0
DB: 9 Gaps: 0

US-09-017-715A-2 (1-127) x BC014098 (1-758)

QY 1 MetAaPvAlPhelYelYsGlyPheSerIleAlaLYsGlyValAlaGlu 20
Db 71 ATGAGTGTCTTCAAGAGGCTTCTCCATCGCCAAAGAGGCGTGTGGTGGTGGAA 130

QY 21 LysThrLYSGInGlyValThrGluAlaAlaGluLYsThrLYSGInGlyValMetYrVal 40
Db 111 AAGACCAAGCAGGCGGTGACGACAGCACTGAGAGACCAAGAGGCGGTCTATGATG 190
QY 41 GJAlaLYsThrLYSGInGlyAsnValAlaGlnSerValThrSerValAlaGluLYsThrLYs 60
Db 131 GGAGCCAAAGACCAAGAGATGTTGTACAGACCGTACCTCAGTGGCCAGAACCAAG 250
QY 61 GJGlnAlaAsnAlaValSerLYsAlaValAlaSerSerValAsnThrValAlaThrLYs 80
Db 251 GAGCAGGCCAACGCGGTGACGAGCTGTGGACACGCTCAACACTGTGGCCACCAAG 310
QY 81 ThrValGlnGluAlaGlnAlaAsnIleAlaValThrSerGlyValAlaArgLYsGluAspLeu 100
Db 311 ACCGTGGAGAGGCGGAGAACATCGCGTCACTCCGGGGTGTGTCGCAAGAGGACTTG 370
QY 101 ArgProSerAlaPProGlnGlnGlnGlyGluAlaSerLYsGlyGluGluValAlaGlu 120
Db 371 AGCCATCTGCCCCCAAGAGAGGAGTGGATCCAAAGAAAGAGAGTGGCAGAG 430
QY 121 GJAlaGlnSerGlyLYsP 127
Db 431 GAGGCCACAGAGTGGGAGAC 451

RESULT 9

AF411524 488 bp mRNA linear PRI 20-SEP-2001
LOCUS Homo sapiens synuclein gamma mRNA, complete cds.
DEFINITION AF411524
ACCESSION AF411524.1 GI:15705904
VERSION
KEYWORDS

SOURCE

ORGANISM Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE 1 (bases 1 to 488)
AUTHORS Han, C., Zhang, B., Peng, X., Yuan, J. and Qiang, B.
TITLE Direct Submission
JOURNAL Submitted (19-AUG-2001) Department of Biochemistry, Institute of
Basic Medical Science, Chinese Academy of Medical Sciences, 5 Dong
Dan San Tiao, Beijing 100005, P.R. China

FEATURES

source
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/db_xref="GI:15705905"
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SAPQEGESKEKEVAERASGSD"

CDS

ORIGIN
Alignment Scores:
Pred. No.: 1,24e-42 Length: 488
Score: 595.00 Matches: 124
Percent Similarity: 99.21% Conservative: 2
Best Local Similarity: 97.64% Mismatches: 1
Query Match: 97.54% Indels: 0
DB: 9 Gaps: 0

US-09-017-715A-2 (1-127) x AF411524 (1-488)

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Db 12 ATGAGTGTCTTCAAGAGGCTTCTCCATCGCCAAAGAGGCGTGTGGTGGTGGAA 71
QY 21 LysThrLYSGInGlyValThrGluAlaAlaGluLYsThrLYSGInGlyValMetYrVal 40

Db	Accession	Length	Score	Percent Similarity	Best Local Similarity	Query Match	Db	Accession	Length	Score	Percent Similarity	Best Local Similarity	Query Match													
Db	72	AAAGCAACAGCAGGGGGTGAACGACAGCTGACAAACCAAGAGGGGTCATGTATGTG	131				Db	132	GGAGCCAAAGACCAAGAGATGTTGTACAGACGTCGACTCATGTGCGCGAGAAACCAAG	191																
Qy	41	GIYALALYSTRLYSGLYASNVAlValGlNserValThrSerValAlaGlulysThrlys	60				Qy	61	GIUGLINALAENVALValISerLYALValValISerSerValAsnThrValAlaThrlys	80																
Db	192	GAGCAGGGCCAAACGCGGTGAGACGAGCTGTGGTGGACACGCTCAACACTGTGGCCACCAAG	251				Db	192	GAGCAGGGCCAAACGCGGTGAGACGAGCTGTGGTGGACACGCTCAACACTGTGGCCACCAAG	251																
Qy	81	ThrValGlUGlUALaGlUASnIleAlaValThrSerGlValValArgLYSGlUASpLeu	100				Qy	252	ACCGTGGAGGAGGCGGAGAAACATCCCGGTCACTCCGGGGTGGTCCGCAAGGAGGACTTG	311																
Db	312	AGGCCATCTGCCCGCCCAACAGAGGGTGAAGCATCAAGAAAGAAAGAGAGTGCACAG	371				Db	312	AGGCCATCTGCCCGCCCAACAGAGGGTGAAGCATCAAGAAAGAAAGAGAGTGCACAG	371																
Qy	121	GIUALaGlNserLYGLYAsp	127				Qy	121	GIUALaGlNserLYGLYAsp	127																
Db	372	GAGGCCAGAGTGGGGGAGAGAC	392				Db	372	GAGGCCAGAGTGGGGGAGAGAC	392																
RESULT 10																										
LOCUS	E36333	720 bp	DNA	linear	PAT 18-JUN-2001																					
DEFINITION	Analytical matter based on synuclein and novel synuclein protein.																									
ACCESSION	E36333																									
VERSION	E36333.1	GI:13022626																								
KEYWORDS	JP 1999239488-A/1.																									
SOURCE	Homo sapiens (human)																									
ORGANISM	Homo sapiens																									
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.																									
AUTHORS	1 (bases 1 to 720)																									
TITLE	Andrew,S.M., Valdimia,R.B. and Arun,M.D.																									
JOURNAL	Analytical matter based on synuclein and novel synuclein protein																									
COMMENT	Patent: JP 1999239488-A 1 07-SEP-1999; THE UNIVERSITY COURT OF THE UNIVERSITY OF ST ANDREWS, NYUROBA LTD OS Homo sapiens (human) PN JP 1999239488-A/1 PD 07-SEP-1999 PF 21-SEP-1998 JP 1998306283 PR 19-SEP-1997 GB 9719879.0 PI ANDREW SMITH MAKKARION, VALDIMIA RUVOVICH I BOCHIMAN, PI ARUN MIAMARD DAVIS PC CI2N15/09,A01K67/027,CI2101/68,G01N33/53,CI2N15/00 CC FH Key Location/Qualifiers FT CDS Location/Qualifiers (49).. (432).																									
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ORIGIN																										
Alignment Scores:																										
Pred. No.: 1.88e-42 Length: 720																										
Score: 595.00 Matches: 124																										
Percent Similarity:																										

OY		41	GlyAlaIyThrLySGluAsnValValGlnSerValThrSerValAlaGluLysThrLys	60
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OY		61	GluGlnAlaAenAlaValSerLysAlaValAlaSerSerValAenThrValAlaThrLys	80
Db		229	GAGCAGGCCAACCGCGTAGCGAGCTGTGTGAGCAGCGTCAACACTGTGTGCCACCAG	288
OY		81	ThrValGluGluValaGluAenIIlaAlaValThrSerGlyValAlaArgLysGluAenLys	100
Db		289	ACCGTGAGGGAGGGAGGAACATCGCGTCACTCCGGGTGTGTCCGCMAAGAGACTTG	348
OY		101	ArgProSerAlaProGlnGlnGluGlyValaSerLysGluLysGluGluValAlaGlu	120
Db		349	AGCCCATGTGCCCCCCCACAGAGAGGCTGTGCATCCAAGAGAAGAAAGAGTGGCACAG	408
OY		121	GluAlaGlnSerGlyGlyASP	127
Db		409	GAGGCCCCAGAGTGGGGGAGAC	429
RESULT 11				
AX004527				
LOCUS	AX004527	720 bp	DNA	linear PAT 24-AUG-2000
DEFINITION	Sequence 1 from Patent EP0908727.			
ACCESSION	AX004527			
VERSION	AX004527.1	GI:9927977		
KEYWORDS				
SOURCE	Homo sapiens (human)			
ORGANISM	Homo sapiens			
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.			
TITLE	1			
JOURNAL	Synuclein-based assay and synuclein protein			
FEATURES	Patent: EP 0908727-A 1 14-APR-1999;			
source	NEUROPA LIMITED (GB); UNIV COURT OF THE UNIVERSITY O (GB)			
CDS	location/Qualifiers			
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ORIGIN				
Alignment Scores:				
Pred. No.:	1.88e-42	Length:	720	
Score:	595.00	Matches:	124	
Percent Similarity:	99.21%	Conservative:	2	
Best Local Similarity:	97.64%	Mismatches:	1	
Query Match:	97.54%	Indels:	0	
DB:	6	Gaps:	0	
US-09-017-715A-2 (1-127) x AX004527 (1-720)				
OY		1	MetaspyralPhelySLySGlyPheSerIIlaAlaLysLysGlyValValGluValaGlu	20
Db		49	ATGATGTCTTCAAGAAAGGCGTTCTCCATGCGCAAGAGGAGCGTGTGGTGGCGGTGAA	108
OY		21	LysThrLysGlnGlyValThrGluAlaAlaGluLysThrLysGluGlyValaMetyVal	40
Db		109	AAGACCAAGCAGGGCGGTGACGGAAGCAGCTGAGAAACCAAGAGGGCGGTGATATGTG	168
OY		41	GlyAlaLysThrLysGluAenValaGlnSerValThrSerValaGluLysThrLys	60
Db		169	GGAGCCAAACCAAGAGAAATGTTGTACAAGGTGACCTCAGTGGCCGAGAACCAAG	228

QY 61 GUGUUAAsnAlaValSerLySaIaValSerSerValAsnThrValaThrLyS 80
 DB 229 GAGCAGGCCAACGCGGTGAGCAGCGTGTGAGCAGCGTCAACACTGTGCGCAACG 288
 QY 81 ThrValAGUUAAGUAsnIleAlaValThrSerGlyValValArgLySGluAspLeu 100
 DB 289 ACCGTGAGAGAGCGGAGAACATCGCGTCACTCCGGGGTGTGGCAAGAGACTTG 348
 QY 101 ArgProSerAlaProGlnGlnGlnGlyGluAlaSerLySGluLySGluGluValaGlu 120
 DB 349 AGCCCATCTGCCCCCACAAGAGAGGCTGTGCATCCAAAGAGAAAGAGAGTGGCAGG 408
 QY 121 GUUaGlnSerGlyGlyAsp 127
 DB 409 GAGGCCCAAGTGGGGAGAC 429

RESULT 12
 AF017256 720 bp mRNA linear PRI 23-SEP-1998
 LOCUS Homo sapiens peryn mRNA, complete cds.
 DEFINITION AF017256
 ACCESSION AF017256.1 GI:3642774
 VERSION AF017256.1 GI:3642774
 KEYWORDS
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 720)
 AUTHORS Ninkina,N.N., Alimova-Kost,M.V., Paterson,J.W., Delaney,L.,
 Cohen,B.B., Imreh,S., Gnuchev,N.V., Davies,A.M. and Buchman,V.L.
 TITLE Organization, expression and polymorphism of the human peryn gene
 JOURNAL Hum. Mol. Genet. 7 (9), 1417-1424 (1998)
 MEDLINE 98367030
 PUBMED 9700196
 REFERENCE 2 (bases 1 to 720)
 AUTHORS Buchman,V.L.
 TITLE Direct Submission
 JOURNAL Submitted (04-AUG-1997) School of Biomedical sciences, Univ. of St.
 Andrews, Bute Medical Buildings, St. Andrews, Fife KY16 9TS,
 Scotland

FEATURES
 source Location/Qualifiers
 1..720
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 49..432
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 /protein_id="AAC36550.1"
 /db_xref="GI:3642775"
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 433..720
 684..689
 706

3'UTR
 polyA_signal
 polyA_site
 706

ORIGIN
 Alignment Scores:
 Pred. No.: 1,886-42 Length: 720
 Score: 595.00 Matches: 124
 Percent Similarity: 99.21% Conservative: 2
 Best Local Similarity: 97.64% Mismatches: 1
 Query Match: 97.54% Indels: 0
 Gaps: 0
 US-09-017-715A-2 (1-127) x AF017256 (1-720)

QY 1 MetaspVAlPhelysLySGlyPheSerIleAlaLySGlyValValaGluValaGlu 20

DB 49 ANGATGTCCTTCAAGAGGGCTTCCATCCGCAAGAGAGGGGTGTGGTCCGATGGA 108
 QY 21 LysThrLySGlnGlyValThrGluAlaGluLyThrLySGlnGlyValMetLyTrVal 40
 DB 109 AAGACCAAGCAGGGGGGTGACCGAACAGCTGTGAAGACCAAGAGGGGTCTGTATGTG 168
 QY 41 GUAAlaLyThrLySGluAsnValaGlnSerValThrSerValaGluLyThrLyS 60
 DB 169 GAGCCAGACCAAGAGAGATTTTACAGACCTGACTCTGAGCCCAAGAACCAAG 228
 QY 61 GUGUUAAsnAlaValSerLySaIaValSerSerValAsnThrValaThrLyS 80
 DB 229 GAGCAGGCCAACGCGGTGAGCAGCGTGTGAGCAGCGTCAACACTGTGCGCAACG 288
 QY 81 ThrValAGUUAAGUAsnIleAlaValThrSerGlyValValArgLySGluAspLeu 100
 DB 289 ACCGTGAGAGAGCGGAGAACATCGCGTCACTCCGGGGTGTGGCAAGAGACTTG 348
 QY 101 ArgProSerAlaProGlnGlnGlnGlyGluAlaSerLySGluLySGluGluValaGlu 120
 DB 349 AGCCCATCTGCCCCCACAAGAGAGGCTGTGCATCCAAAGAGAAAGAGAGTGGCAGG 408
 QY 121 GUUaGlnSerGlyGlyAsp 127
 DB 409 GAGGCCCAAGTGGGGAGAC 429

RESULT 13
 BV177827 738 bp DNA linear STS 10-JUN-2004
 LOCUS sqm97020 Human DNA (Sequenom) Homo sapiens STS genomic, sequence
 DEFINITION BV177827
 ACCESSION BV177827.1 GI:48014020
 VERSION BV177827.1 GI:48014020
 KEYWORDS
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 738)
 AUTHORS Nelson,R.M., Warrnellos,G., Kammerer,S., Hoyal,C.R., Shi,M.M.,
 Cantor,C.R. and Braun,A.
 TITLE Large-Scale Validation of Single Nucleotide Polymorphisms in Gene
 JOURNAL Regions
 Genome Res. (2004) In press
 COMMENT
 Contact: Andreas Braun
 Pharmaceuticals division
 Sequenom, Inc.
 3595 John Hopkins Court, San Diego, CA 92121, USA
 Tel: 18582029018
 Fax: 18582029020
 Email: abraun@sequenom.com
 Primer A: No primer sequence submitted
 Primer B: No primer sequence submitted
 STS size: 738.

FEATURES
 source Location/Qualifiers
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 /mol_type="genomic DNA"
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ORIGIN
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 Alignment Scores:
 Pred. No.: 4,486-37 Length: 738
 Score: 532.50 Matches: 120
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 Best Local Similarity: 94.49% Mismatches: 5
 Query Match: 87.30% Indels: 2
 Gaps: 0
 DB: 11

US-09-017-715A-2 (1-127) x BV177827 (1-738)

QY 1 MetaspValpbelvlysglypheserlleaAlalyserglyValaIGlyAlaValaIGlu 20
Db 53 ATGGATGCTTCAAGAAAGGGCTCTCCATCCCAAGAGGGCGTGGTGGCGGTGAA 112
QY 21 LyethrlysgInglyValThrgluAlaAGluThlyrThlyrsgluValMetTyVal 40
Db 113 TAGCAACCAAGCTGTCTTACGAAGCACTGAGAACCAAGAGGGGTG-ATGTATGT- 170
QY 41 GYAlalyThrlysgluAenValaIGlnserValThrserValaIGluThlyr 60
Db 171 GAGACCAAGCAAGCAAGAAATGTGTACAGAGCTGACCTCAGTGGCCGAGAACCAAG 230
QY 61 GlnGlnAlaAenAlaValserlyAlaValaIGlnserValaenThrValaThlyr 80
Db 231 GAGAGGCAACCAAGCGGTGAGAGGCTGTGTGAGACCGTCAACACTGTGGCCACCAAG 290
QY 81 ThrValIGluAlaIGluAenValaValaThrserlyValaArglysgluApleu 100
Db 291 ACCGTGAGAGAGCGGAGAACATCGCGTCACTCCGGGTGTGTCCAGAGAGGACTTG 350
QY 101 ArgProserAlaProGInglyValaIGluAserlysgluValaIGluValaIGlu 120
Db 351 AGGCACTGTGCCCCCAACAGAGGCTGCGCATCCAAAGAAAGAGAGTGGCAGAG 410
QY 121 GluAlaGlnserglyValasp 127
Db 411 GAGGCCCAAGAGTGGGGAGAC 431

RESULT 14

AF219257 677 bp mRNA linear MAM 09-FEB-2000
LOCUS Bos taurus synoretin mRNA, complete cds.
DEFINITION
AF219257
AF219257.1 GI:6942173

VERSION
KEYWORDS
SOURCE
ORGANISM

Bos taurus (cow)

Bos taurus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
Bovinae; Bos.

REFERENCE
1 (bases 1 to 677)
Surguchov, A., Surgucheva, I., Solesio, E. and Baehr, W.
Synoretin-A new protein belonging to the synuclein family
Mol. Cell. Neurosci. 13 (2), 95-103 (1999)

AUTHORS
TITLE
JOURNAL
MEDLINE
PUBMED
2 (bases 1 to 677)
Surguchov, A., Surgucheva, I., Baehr, W. and Solesio, E.
Direct Submission
Submitted (27-DEC-1999) Ophthalmology, Washington University School
of Medicine in St. Louis, 660 South Euclid, St. Louis, MO 63108,
USA

FEATURES

source

1. 677
/organism="Bos taurus"
/mol_type="mRNA"
/db_xref="taxon:9913"

1. 384
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/protein_id="AAF2342.1"
/db_xref="GI:6942174"

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ORIGIN

Alignment Scores:
Pred. No.:

2.67e-36 Length: 677

Score: 523.00 Matches: 109
Percent Similarity: 91.34% Conservative: 7
Best Local Similarity: 85.83% Mismatches: 11
Query Match: 85.74% Indels: 0
DB: 4 Gaps: 0

US-09-017-715A-2 (1-127) x AF219257 (1-677)

QY 1 MetaspValpbelvlysglypheserlleaAlalyserglyValaIGlyAlaValaIGlu 20
Db 1 ATGGACCTCTTCAAGAAAGGGCTCTCCATTCAGAGAGGGGTGGTGGCGGTGAA 60
QY 21 LyethrlysgInglyValThrgluAlaAGluThlyrThlyrsgluValMetTyVal 40
Db 61 AAGCAACCAAGGAGTGTGACAGAGCACTGAGAACCAAGAGGGGTGTATATGTG 120
QY 41 GYAlalyThrlysgluAenValaIGlnserValThrserValaIGluThlyr 60
Db 121 GAGCTAAGCAACCAAGAGGGGTGTGTGACAGTGTGACTTCACTGCTGAGAACCAAG 180
QY 61 GlnGlnAlaAenAlaValserlyAlaValaIGlnserValaenThrValaThlyr 80
Db 181 GAGCAAGCCCAAGCGGTGAGAGGCGGTGTCTCCAGTGTCAACACTGTGGCCACCAAG 240
QY 81 ThrValIGluAlaIGluAenValaValaThrserlyValaArglysgluApleu 100
Db 241 ACTGTGAGAGAGGTGTGAGAACATTCAGTCACTCTGAGTGTGTCACAGAGAGGCCCTG 300
QY 101 ArgProserAlaProGInglyValaIGluAserlysgluValaIGluValaIGlu 120
Db 301 AAGCAACCTGTCCCTTCAAGAGAGGATGAGAGCAAGCAAGAGAGAGGCGCTTG 360
QY 121 GluAlaGlnserglyValasp 127
Db 361 GAGACCAAGAGTGGGGAGAT 381

RESULT 15

E36334 727 bp DNA linear PAT 18-JUN-2001
LOCUS Analytical matter based on synuclein and novel synuclein protein.
DEFINITION
E36334
E36334.1 GI:13022627
VERSION
E36334.1 GI:13022627
KEYWORDS
JP 1999239488-A/2.
SOURCE
Mus musculus (house mouse)

ORGANISM
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE
1 (bases 1 to 727)
Andrew, S.M., Valdimia, R.B. and Arun, M.D.
Analytical matter based on synuclein and novel synuclein protein
Patent: JP 1999239488-A 2 07-SEP-1999;
THE UNIVERSITY COURT OF THE UNIVERSITY OF ST ANDREWS, NYUROBA LTD

AUTHORS
TITLE
JOURNAL
COMMENT
OS Mus musculus (mouse)
PN JP 1999239488-A/2
PD 07-SEP-1999
PF 21-SEP-1998 JP 1998306283
PR 19-SEP-1997 GB 9719879.0
PI ANDREW SMITH MAKARION, VALDIMIA RUVOVICH BUCHIMAN, PI ARUN
MILNARD DAVIS

PC C12N15/09, A01K67/027, C12Q1/68, G01N33/53, C12N15/00 CC
FH Key
FT CDS Location/Qualifiers (69)..(440).

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1. 727
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/db_xref="taxon:10090"

1. 727
Location/Qualifiers

1. 727
Location/Qualifiers

1. 727
Location/Qualifiers

ORIGIN

Alignment Scores:
Pred. No.:

4.93e-34 Length: 727
Score: 497.00 Matches: 108
Percent Similarity: 89.76% Conservative: 6

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - nucleic search, using frame_plus_p2n model

Run on: May 4, 2005, 10:17:21 ; Search time 65.0738 Seconds

(without alignments) 748,404 Million cell updates/sec

Title: US-09-017-715A-2_COPY_120_127

Perfect score: 41

Sequence: 1 EEAQSGSD 8

Scoring table:

BLOSUM62

Xgapop 10.0 , Ygapext 0.5

Ygapop 10.0 , Ygapext 0.5

Fgapop 6.0 , Fgapext 7.0

Delop 6.0 , Delext 7.0

Searched: 5642217 seqs, 3043843248 residues

Total number of hits satisfying chosen parameters: 11284434

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Command line parameters:

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-Q=/cg2_1/USPTO.spool_h/US09017715/runat_04052005_100747_25719/app_query.fasta_1.661

-DB=Published Applications NA -QFMT=fastab -SUFFIX=emb -MINMATCH=0.1

-LOOPCL=0 -LOOPEXT=0 -UNITS=bases -START=1 -END=1 -MATRIX=blone62

-TRANS=human40.cct -LIST=45 -DOCLIGN=200 -THR SCORE=pct -THR MAX=100

-THR_MIN=0 -ALIGN=15 -MODE=LOCAL -OUTFMT=ptc -NORM=ext -HEAPSIZE=500 -MINLEN=0

-MAXLEN=2000000000 -USER=US09017715 -@CGN_1_1_684 -@runat_04052005_100747_25719

-NCPUPU=6 -ICPU=3 -NO MMAP -LARGQUERY -NEG SCORES=0 -WAIT -DSBLOCK=100

-LONGLOG -DEV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5

-FGAPOP=6 -FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

Published Applications NA:*

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2: /cg2_6/ptodata/1/pubpna/PCT_NEW_PUB.seq:*

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12: /cg2_6/ptodata/1/pubpna/US09_NEW_PUB.seq:*

13: /cg2_6/ptodata/1/pubpna/US10A_PUBCOMB.seq:*

14: /cg2_6/ptodata/1/pubpna/US10B_PUBCOMB.seq:*

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19: /cg2_6/ptodata/1/pubpna/US10_NEW_PUB.seq:*

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1 41 100.0 384 19 US-10-826-157-5 Sequence 5, Appl1

2 41 100.0 478 9 US-09-925-298-172 Sequence 172, App

3 41 100.0 478 14 US-10-102-806-172 Sequence 172, App

4 41 100.0 478 10 US-09-918-995-2705 Sequence 2705, App

5 41 100.0 550 9 US-09-954-531-613 Sequence 613, App

6 41 100.0 550 17 US-10-453-478-1 Sequence 1, Appl1

7 41 100.0 550 17 US-10-843-641A-1680 Sequence 1680, App

8 41 100.0 720 14 US-10-097-340-297 Sequence 297, App

9 41 100.0 720 17 US-10-282-174-469 Sequence 469, App

10 41 100.0 720 9 US-10-600-009-469 Sequence 469, App

11 41 100.0 796 9 US-09-925-298-171 Sequence 171, App

12 41 100.0 796 14 US-10-102-806-171 Sequence 171, App

13 38 92.7 1125 17 US-10-369-493-33286 Sequence 33286, A

14 37 90.2 442 18 US-10-425-115-90466 Sequence 90466, A

15 37 90.2 2185 18 US-10-437-963-8217 Sequence 8217, App

16 37 90.2 106236 19 US-10-741-600-17759 Sequence 17759, A

17 36 87.8 357 18 US-10-437-963-43087 Sequence 43087, A

18 36 87.8 406 17 US-10-424-599-15323 Sequence 15323, A

19 36 87.8 1174 13 US-10-027-632-123222 Sequence 123222, A

20 36 87.8 1174 13 US-10-027-632-123222 Sequence 123223, A

21 36 87.8 1174 17 US-10-027-632-123222 Sequence 123223, A

22 36 87.8 1174 17 US-10-027-632-123223 Sequence 123223, A

23 36 87.8 2116 17 US-10-104-047-1099 Sequence 1099, App

24 36 87.8 3016 18 US-10-602-494-48 Sequence 48, Appl

25 36 87.8 4985 13 US-10-071-223-1 Sequence 1, Appl1

26 36 87.8 4985 19 US-10-896-552-1 Sequence 187, App

27 36 87.8 5372 9 US-09-751-1008-98 Sequence 98, Appl

28 36 87.8 5515 18 US-10-470-565-1 Sequence 1, Appl1

29 36 87.8 225666 18 US-10-425-115-65435 Sequence 65435, A

30 35 85.4 291 18 US-10-767-701-27423 Sequence 27423, A

31 35 85.4 397 18 US-10-021-323-3503 Sequence 3503, App

32 35 85.4 547 16 US-10-029-386-6547 Sequence 6547, App

33 35 85.4 775 17 US-10-425-114-6521 Sequence 6521, App

34 35 85.4 588 18 US-10-437-963-6679 Sequence 6679, App

35 35 85.4 775 17 US-10-425-115-11974 Sequence 11974, A

36 35 85.4 884 18 US-10-425-115-119416 Sequence 119416, A

37 35 85.4 918 17 US-10-369-493-44914 Sequence 44914, A

38 35 85.4 1901 17 US-10-424-599-131648 Sequence 131648, A

39 35 85.4 1928 17 US-10-425-114-24113 Sequence 24113, A

40 35 85.4 2047 17 US-10-425-114-27013 Sequence 27013, A

ALIGNMENTS

US-10-826-157-5

Sequence 5, Application US/10826157

Publication No. US20050064548A1

GENERAL INFORMATION:

APPLICANT: Lindquist, Susan L.

APPLICANT: Oliveira, Tiago

TITLE OF INVENTION: YEAST ECTOPOLOGICALLY EXPRESSING ABNORMALLY

FILE REFERENCE: 17481-003001

CURRENT APPLICATION NUMBER: US/10/826,157

CURRENT FILING DATE: 2004-04-16

PRIOR APPLICATION NUMBER: US 60/472,317

PRIOR FILING DATE: 2003-05-20

PRIOR APPLICATION NUMBER: US 60/463,284

PRIOR FILING DATE: 2003-04-16

NUMBER OF SEQ ID NOS: 8

SOFTWARE: FastSeq for Windows Version 4.0

SEQ ID NO 5

LENGTH: 384

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-826-157-5

Alignment Scores:
Pred. No.: 20.2      Length: 384
Score: 41.00      Matches: 8
Percent Similarity: 100.00%      Conservative: 0
Best Local Similarity: 100.00%      Mismatches: 0
Query Match: 100.00%      Indels: 0
DB: 19      Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x US-10-826-157-5 (1-384)
QY 1 GIUGUAlaGInSerGlyGlyasp 8
DB 358 GAGGAGGCCAGAGTGGGGAGAC 381

RESULT 2
US-09-925-298-172
; Sequence 172, Application US/09925298
; Publication No. US20020039764A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins and Antibodies
; FILE REFERENCE: PA103
; CURRENT APPLICATION NUMBER: US/09/925,298
; CURRENT FILING DATE: 2001-08-10
; PRIOR APPLICATION NUMBER: PCT/US00/05881
; PRIOR FILING DATE: 2000-03-08
; PRIOR APPLICATION NUMBER: 60/124,270
; PRIOR FILING DATE: 1999-03-12
; NUMBER OF SEQ ID NOS: 846
; SOFTWARE: Patentln Ver. 2.0
; SEQ ID NO 172
; LENGTH: 478
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-925-298-172

Alignment Scores:
Pred. No.: 24.4      Length: 478
Score: 41.00      Matches: 8
Percent Similarity: 100.00%      Conservative: 0
Best Local Similarity: 100.00%      Mismatches: 0
Query Match: 100.00%      Indels: 0
DB: 9      Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x US-09-925-298-172 (1-478)
QY 1 GIUGUAlaGInSerGlyGlyasp 8
DB 149 GAGGAGGCCAGAGTGGGGAGAC 172

RESULT 3
US-10-102-806-172
; Sequence 172, Application US/10102806
; Publication No. US20030054421A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins and Antibodies
; FILE REFERENCE: PA103P1C1
; CURRENT APPLICATION NUMBER: US/10/102,806
; CURRENT FILING DATE: 2002-03-22
; PRIOR APPLICATION NUMBER: 09/925,298
; PRIOR FILING DATE: 2001-08-10
; PRIOR APPLICATION NUMBER: PCT/US00/05881
; PRIOR FILING DATE: 2000-03-08
; PRIOR APPLICATION NUMBER: 60/124,270
; PRIOR FILING DATE: 1999-03-12
; NUMBER OF SEQ ID NOS: 846
; SOFTWARE: Patentln Ver. 2.0
; SEQ ID NO 172
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; LENGTH: 478
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-102-806-172

Alignment Scores:
Pred. No.: 24.4      Length: 478
Score: 41.00      Matches: 8
Percent Similarity: 100.00%      Conservative: 0
Best Local Similarity: 100.00%      Mismatches: 0
Query Match: 100.00%      Indels: 0
DB: 14      Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x US-10-102-806-172 (1-478)
QY 1 GIUGUAlaGInSerGlyGlyasp 8
DB 149 GAGGAGGCCAGAGTGGGGAGAC 172

RESULT 4
US-09-918-995-2705
; Sequence 2705, Application US/09918995
; Publication No. US20030073623A1
; GENERAL INFORMATION:
; APPLICANT: Hyseq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
; FROM VARIOUS CDNA LIBRARIES
; FILE REFERENCE: 20411-756
; CURRENT APPLICATION NUMBER: US/09/918,995
; CURRENT FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: US/09/235,076
; PRIOR FILING DATE: 1999-01-20
; NUMBER OF SEQ ID NOS: 38054
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 2705
; LENGTH: 479
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(479)
; OTHER INFORMATION: n = A,T,C or G
US-09-918-995-2705

Alignment Scores:
Pred. No.: 24.4      Length: 479
Score: 41.00      Matches: 8
Percent Similarity: 100.00%      Conservative: 0
Best Local Similarity: 100.00%      Mismatches: 0
Query Match: 100.00%      Indels: 0
DB: 10      Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x US-09-918-995-2705 (1-479)
QY 1 GIUGUAlaGInSerGlyGlyasp 8
DB 371 GAGGAGGCCAGAGTGGGGAGAC 394

RESULT 5
US-09-954-531-613
; Sequence 613, Application US/09954531
; Patent No. US20020165180A1
; GENERAL INFORMATION:
; APPLICANT: Weaver, Zoe
; TITLE OF INVENTION: Process for Identifying Anti-Cancer Therapeutic Agents Using Cance
; FILE REFERENCE: 689290-77
; CURRENT APPLICATION NUMBER: US/09/954,531
; CURRENT FILING DATE: 2002-05-02
; PRIOR APPLICATION NUMBER: US/60/233,133
; PRIOR FILING DATE: 2000-09-18
; PRIOR APPLICATION NUMBER: US/60/234,009
; PRIOR FILING DATE: 2000-09-20
```



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PRIOR APPLICATION NUMBER: US/60/234,034
PRIOR FILING DATE: 2000-09-20
PRIOR APPLICATION NUMBER: US/60/234,509
PRIOR FILING DATE: 2000-09-22
PRIOR APPLICATION NUMBER: US/60/234,567
PRIOR FILING DATE: 2000-09-22
NUMBER OF SEQ ID NOS: 1392
SOFTWARE: PatentIn version 3.0
SEQ ID NO: 613
LENGTH: 550
TYPE: DNA
ORGANISM: Homo sapiens
US-09-954-531-613

Alignment Scores:
Pred. No.: 27.5      Length: 550
Score: 41.00        Matches: 8
Percent Similarity: 100.00%    Conservative: 0
Best Local Similarity: 100.00%    Mismatches: 0
Query Match: 100.00%    Indels: 0
DB: 9      Gaps: 0

US-09-017-715a-2_COPY_120_127 (1-8) x US-09-954-531-613 (1-550)

Qy 1 GluGluAlaGlnSerGlyGlyAsp 8
Db 369 GAGGAGGCCGACGACTGGGGGAGAC 392

RESULT 6
US-10-453-478-1
; Sequence 1, Application US/10453478
; Publication No. US20030208043A1
GENERAL INFORMATION:
APPLICANT: Paul Moore, Reiner Gentz, Hongjin Ji,
Jian Ni and Jing-Shan Hu
TITLE OF INVENTION: Human Genes, Sequences and
Expression Products
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: CAEHLA, BYRNE, BAIN, GILFILLAN,
CROCHT, STEWART & OLSTEIN
STREET: 6 BECKER FARM ROAD
CITY: ROSELAND
STATE: NEW JERSEY
COUNTRY: USA
ZIP: 07068
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5 INCH DISKETTE
COMPUTER: IBM PS/2
OPERATING SYSTEM: MS-DOS
SOFTWARE: WORD PERFECT 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/10/453,478
FILING DATE: 04-Jun-2003
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/08/705,771
FILING DATE: August 30, 1996
ATTORNEY/AGENT INFORMATION:
NAME: MULHINS, J.G.
REGISTRATION NUMBER: 33,073
REFERENCE/DOCKET NUMBER: 325800-346 (PFI96)
TELECOMMUNICATION INFORMATION:
TELEPHONE: 973-994-1700
TELEFAX: 973-994-1744
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 550 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
SEQUENCE DESCRIPTION: SEQ ID NO: 1:

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US-10-453-478-1

Alignment Scores:
Pred. No.: 27.5      Length: 550
Score: 41.00        Matches: 8
Percent Similarity: 100.00%    Conservative: 0
Best Local Similarity: 100.00%    Mismatches: 0
Query Match: 100.00%    Indels: 0
DB: 17      Gaps: 0

US-09-017-715a-2_COPY_120_127 (1-8) x US-10-453-478-1 (1-550)

Qy 1 GluGluAlaGlnSerGlyGlyAsp 8
Db 369 GAGGAGGCCGACGACTGGGGGAGAC 392

RESULT 7
US-10-843-641A-1680
; Sequence 1680, Application US/10843641A
; Publication No. US20050064454A1
GENERAL INFORMATION:
APPLICANT: Avalon Pharmaceuticals, Inc.
TITLE OF INVENTION: Cancer Gene Determination and Therapeutic Screening Using
FILE REFERENCE: 689290-189
CURRENT APPLICATION NUMBER: US/10/843,641A
CURRENT FILING DATE: 2004-05-12
PRIOR APPLICATION NUMBER: US/09/873,367
PRIOR FILING DATE: 2001-06-05
PRIOR APPLICATION NUMBER: US/09/954,531
PRIOR FILING DATE: 2001-09-18
PRIOR APPLICATION NUMBER: US/09/954,456
PRIOR FILING DATE: 2001-09-25
PRIOR APPLICATION NUMBER: US/09/962,436
PRIOR FILING DATE: 2001-09-25
PRIOR APPLICATION NUMBER: US/09/962,832
PRIOR FILING DATE: 2001-09-25
PRIOR APPLICATION NUMBER: US/09/964,824
PRIOR FILING DATE: 2001-09-27
PRIOR APPLICATION NUMBER: US/09/967,768
PRIOR FILING DATE: 2001-09-28
PRIOR APPLICATION NUMBER: US/09/968,007
PRIOR FILING DATE: 2001-10-02
PRIOR APPLICATION NUMBER: US/09/969,347
PRIOR FILING DATE: 2001-10-02
PRIOR APPLICATION NUMBER: US/09/969,708
PRIOR FILING DATE: 2001-10-03
Remaining Prior Application data removed - See File Wrapper or PALM.
NUMBER OF SEQ ID NOS: 8447
SOFTWARE: PatentIn version 3.0
SEQ ID NO: 1680
LENGTH: 550
TYPE: DNA
ORGANISM: Homo sapiens
US-10-843-641A-1680

Alignment Scores:
Pred. No.: 27.5      Length: 550
Score: 41.00        Matches: 8
Percent Similarity: 100.00%    Conservative: 0
Best Local Similarity: 100.00%    Mismatches: 0
Query Match: 100.00%    Indels: 0
DB: 19      Gaps: 0

US-09-017-715a-2_COPY_120_127 (1-8) x US-10-843-641A-1680 (1-550)

Qy 1 GluGluAlaGlnSerGlyGlyAsp 8
Db 369 GAGGAGGCCGACGACTGGGGGAGAC 392

RESULT 8
US-10-097-340-297
; Sequence 297, Application US/10097340

```

```
Publication No. US20030087250A1
GENERAL INFORMATION:
APPLICANT: John MONAHAN
APPLICANT: Manjula GANNANARAPU
APPLICANT: Sebastian HOERSCH
APPLICANT: Shubhangi KAMATKAR
APPLICANT: Steve G. KOVATS
APPLICANT: Rachel E. MEYERS
APPLICANT: Michael MORRISSEY
APPLICANT: Peter OLANDT
APPLICANT: Ami SEN
APPLICANT: Peter VEIBY
APPLICANT: Gordon B. MILLS
APPLICANT: Robert C. BAST, Jr.
APPLICANT: Karen LU
APPLICANT: Rosemarie SCHMANDT
APPLICANT: Xumei ZHAO
APPLICANT: Karen GIATT
TITLE OF INVENTION: Nucleic Acid Molecules and Proteins For The Identification,
FILE REFERENCE: MRI-030
CURRENT APPLICATION NUMBER: US/10/097,340
CURRENT FILING DATE: 2002-03-14
PRIOR APPLICATION NUMBER: 60/276,025
PRIOR FILING DATE: 2001-03-14
PRIOR APPLICATION NUMBER: 60/325,149
PRIOR FILING DATE: 2001-09-26
PRIOR APPLICATION NUMBER: 60/276,026
PRIOR FILING DATE: 2001-03-14
PRIOR APPLICATION NUMBER: 60/324,967
PRIOR FILING DATE: 2001/09/26
PRIOR APPLICATION NUMBER: 60/311,732
PRIOR FILING DATE: 2001-08-10
PRIOR APPLICATION NUMBER: 60/325,102
PRIOR FILING DATE: 2001-09-26
PRIOR APPLICATION NUMBER: 60/323,580
PRIOR FILING DATE: 2001-09-19
NUMBER OF SEQ ID NOS: 363
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 297
LENGTH: 720
TYPE: DNA
ORGANISM: Homo sapiens
US-10-097-340-297

Alignment Scores:
Pred. No.: 34.8      Length: 720
Score: 41.00      Matches: 8
Percent Similarity: 100.00%      Conservative: 0
Best Local Similarity: 100.00%      Mismatches: 0
Query Match: 100.00%      Indels: 0
DB: 14      Gaps: 0

US-09-017-715a-2_copy_120_127 (1-8) x US-10-097-340-297 (1-720)
Qy 1 GIUGUAAGInSerGIyGlyasp 8
Db 406 GAGGAGCCCAAGAGTGGGGAGAC 429

RESULT 9
US-10-282-174-469
Sequence 469, Application US/10282174
Publication No. US20030224380A1
GENERAL INFORMATION:
APPLICANT: Becker, Kenneth David
APPLICANT: Velicelebi, Gonul
APPLICANT: Eliot, Kathryn J.
APPLICANT: Wang, Xin
APPLICANT: Tanzi, Rudolph E.
APPLICANT: Bertam, Lars
APPLICANT: Saunders, Aleister J.
APPLICANT: Mullin, Kristina M.
APPLICANT: Sampson, Andrew Johnson
```

```
APPLICANT: Blacker, Deborah Lynne
TITLE OF INVENTION: GENES AND POLYMORPHISMS ON CHROMOSOME 10
TITLE OF INVENTION: ASSOCIATED WITH ALZHEIMER'S DISEASE AND OTHER
TITLE OF INVENTION: NEURODEGENERATIVE DISEASES
FILE REFERENCE: 37481-3308
CURRENT APPLICATION NUMBER: US/10/282,174
CURRENT FILING DATE: 2002-10-25
PRIOR APPLICATION NUMBER: US 60/339,525
PRIOR FILING DATE: 2001-10-25
PRIOR APPLICATION NUMBER: US 60/338,010
PRIOR FILING DATE: 2001-11-08
PRIOR APPLICATION NUMBER: US 60/336,929
PRIOR FILING DATE: 2001-11-08
PRIOR APPLICATION NUMBER: US 60/337,052
PRIOR FILING DATE: 2001-12-04
PRIOR APPLICATION NUMBER: US 60/368,919
PRIOR FILING DATE: 2002-03-28
NUMBER OF SEQ ID NOS: 564
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 469
LENGTH: 720
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: allele
LOCATION: 30,57,85,243,250,377,512,531,555,561,672
OTHER INFORMATION: N is any
US-10-282-174-469

Alignment Scores:
Pred. No.: 34.8      Length: 720
Score: 41.00      Matches: 8
Percent Similarity: 100.00%      Conservative: 0
Best Local Similarity: 100.00%      Mismatches: 0
Query Match: 100.00%      Indels: 0
DB: 17      Gaps: 0

US-09-017-715a-2_copy_120_127 (1-8) x US-10-282-174-469 (1-720)
Qy 1 GIUGUAAGInSerGIyGlyasp 8
Db 406 GAGGAGCCCAAGAGTGGGGAGAC 429

RESULT 10
US-10-600-009-469
Sequence 469, Application US/10600009
Publication No. US2005009031A1
GENERAL INFORMATION:
APPLICANT: Becker, Kenneth David
APPLICANT: Velicelebi, Gonul
APPLICANT: Eliot, Kathryn J.
APPLICANT: Wang, Xin
APPLICANT: Tanzi, Rudolph E.
APPLICANT: Bertam, Lars
APPLICANT: Saunders, Aleister J.
APPLICANT: Mullin, Kristina M.
APPLICANT: Sampson, Andrew Johnson
APPLICANT: Blacker, Deborah Lynne
TITLE OF INVENTION: GENES AND POLYMORPHISMS ON CHROMOSOME 10
TITLE OF INVENTION: ASSOCIATED WITH ALZHEIMER'S DISEASE AND OTHER
TITLE OF INVENTION: NEURODEGENERATIVE DISEASES
FILE REFERENCE: 37481-3308B
CURRENT APPLICATION NUMBER: US/10/600,009
CURRENT FILING DATE: 2003-06-18
PRIOR APPLICATION NUMBER: US 60/339,525
PRIOR FILING DATE: 2001-10-25
PRIOR APPLICATION NUMBER: US 60/338,010
PRIOR FILING DATE: 2001-11-08
PRIOR APPLICATION NUMBER: US 60/336,929
PRIOR FILING DATE: 2001-11-08
PRIOR APPLICATION NUMBER: US 60/337,052
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; PRIOR FILING DATE: 2001-11-09
; PRIOR APPLICATION NUMBER: US 60/337,052
; PRIOR FILING DATE: 2001-12-04
; PRIOR APPLICATION NUMBER: US 60/368,919
; PRIOR FILING DATE: 2002-03-28
; PRIOR APPLICATION NUMBER: US 10/282,174
; PRIOR FILING DATE: 2002-10-25
; NUMBER OF SEQ ID NOS: 564
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 469
; LENGTH: 720
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: allele
; LOCATION: 30,57,85,243,250,377,512,531,555,561,672
; OTHER INFORMATION: N is any
; US-10-600-009-469

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```

Alignment Scores:
Pred. No.: 34.8      Length: 720
Score: 41.00      Matches: 8
Percent Similarity: 100.00%      Conservative: 0
Best Local Similarity: 100.00%      Mismatches: 0
Query Match: 100.00%      Indels: 0
DB: 19      Gaps: 0

```

US-09-017-715A-2_COPY_120_127 (1-8) x US-10-600-009-469 (1-720)

OY 1 GluGluaIaGInserGIyGIyAap 8

Db 406 GAGGAGGCCAGAGTGGGAGGAC 429

```

RESULT 11
US-09-925-298-171
; Sequence 171, Application US/09925298
; Publication No. US20020039764A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins and Antibodies
; FILE REFERENCE: PA103
; CURRENT APPLICATION NUMBER: US/09/925,298
; CURRENT FILING DATE: 2001-08-10
; PRIOR APPLICATION NUMBER: PCT/US00/05881
; PRIOR FILING DATE: 2000-03-08
; PRIOR APPLICATION NUMBER: 60/124,270
; PRIOR FILING DATE: 1999-03-12
; NUMBER OF SEQ ID NOS: 846
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 171
; LENGTH: 796
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-925-298-171

```

```

Alignment Scores:
Pred. No.: 37.9      Length: 796
Score: 41.00      Matches: 8
Percent Similarity: 100.00%      Conservative: 0
Best Local Similarity: 100.00%      Mismatches: 0
Query Match: 100.00%      Indels: 0
DB: 9      Gaps: 0

```

US-09-017-715A-2_COPY_120_127 (1-8) x US-09-925-298-171 (1-796)

OY 1 GluGluaIaGInserGIyGIyAap 8

Db 466 GAGGAGGCCAGAGTGGGAGGAC 489

```

RESULT 12
US-10-102-806-171
; Sequence 171, Application US/10102806
; Publication No. US20030054421A1

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```

; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins and Antibodies
; FILE REFERENCE: PA103P1C1
; CURRENT APPLICATION NUMBER: US/10/102,806
; CURRENT FILING DATE: 2002-03-22
; PRIOR APPLICATION NUMBER: 09/925,298
; PRIOR FILING DATE: 2001-08-10
; PRIOR APPLICATION NUMBER: PCT/US00/05881
; PRIOR FILING DATE: 2000-03-08
; PRIOR APPLICATION NUMBER: 60/124,270
; PRIOR FILING DATE: 1999-03-12
; NUMBER OF SEQ ID NOS: 846
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 171
; LENGTH: 796
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-10-102-806-171

```

```

Alignment Scores:
Pred. No.: 37.9      Length: 796
Score: 41.00      Matches: 8
Percent Similarity: 100.00%      Conservative: 0
Best Local Similarity: 100.00%      Mismatches: 0
Query Match: 100.00%      Indels: 0
DB: 14      Gaps: 0

```

US-09-017-715A-2_COPY_120_127 (1-8) x US-10-102-806-171 (1-796)

OY 1 GluGluaIaGInserGIyGIyAap 8

Db 466 GAGGAGGCCAGAGTGGGAGGAC 489

```

RESULT 13
US-10-369-493-33286/c
; Sequence 33286, Application US/10369493
; Publication No. US30030233675A1
; GENERAL INFORMATION:
; APPLICANT: Cao, Yongwei
; APPLICANT: Hinkle, Gregory J.
; APPLICANT: Slater, Steven C.
; APPLICANT: Goldman, Barry S.
; APPLICANT: Chen, Xianfeng
; TITLE OF INVENTION: EXPRESSION OF MICROBIAL PROTEINS IN PLANTS FOR PRODUCTION OF
; FILE REFERENCE: 38-10(52052)B
; CURRENT APPLICATION NUMBER: US/10/369,493
; CURRENT FILING DATE: 2003-02-28
; PRIOR APPLICATION NUMBER: US 60/360,039
; PRIOR FILING DATE: 2002-02-21
; NUMBER OF SEQ ID NOS: 47374
; SEQ ID NO 33286
; LENGTH: 1125
; TYPE: DNA
; ORGANISM: Desulfotobacterium hafnienae
; US-10-369-493-33286

```

```

Alignment Scores:
Pred. No.: 201      Length: 1125
Score: 38.00      Matches: 7
Percent Similarity: 100.00%      Conservative: 1
Best Local Similarity: 87.50%      Mismatches: 0
Query Match: 92.68%      Indels: 0
DB: 17      Gaps: 0

```

US-09-017-715A-2_COPY_120_127 (1-8) x US-10-369-493-33286 (1-1125)

OY 1 GluGluaIaGInserGIyGIyAap 8

Db 609 GAGGAGGCCAGAGTGGGAGGAC 586

RESULT 14

US-10-425-115-90466/c
; Sequence 90466, Application US/10425115
; Publication No. US20040214272A1
; GENERAL INFORMATION:
; APPLICANT: La Rosa, Thomas J.
; APPLICANT: Kovalic, David K.
; APPLICANT: Zhou, Yihua
; APPLICANT: Cao, Yongwei
; TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated With
; FILE OF INVENTION: Plants
; FILE REFERENCE: 38-21(53222)B
; CURRENT APPLICATION NUMBER: US/10/425,115
; CURRENT FILING DATE: 2003-04-28
; NUMBER OF SEQ ID NOS: 369326
; SEQ ID NO 90466
; LENGTH: 442
; TYPE: DNA
; ORGANISM: Zea mays
; FEATURE:
; NAME/KEY: unsure
; LOCATION: (1)..(442)
; OTHER INFORMATION: unsure at all n locations
; FEATURE:
; OTHER INFORMATION: Clone ID: MRT4577_182507C.1
US-10-425-115-90466

Alignment Scores:
Pred. No.: 142 Length: 442
Score: 37.00 Matches: 7
Percent Similarity: 100.00% Conservative: 1
Best Local Similarity: 87.50% Mismatches: 0
Query Match: 90.24% Indels: 0
DB: 18 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x US-10-425-115-90466 (1-442)

Qy 1 GUGUUAAGInserGlyasp 8
Db 55 GAGGAGCGAGGAGCGGAGGAGC 32

RESULT 15
US-10-437-963-8217
; Sequence 8217, Application US/10437963
; Publication No. US20040123343A1
; GENERAL INFORMATION:
; APPLICANT: La Rosa, Thomas J.
; APPLICANT: Kovalic, David K.
; APPLICANT: Zhou, Yihua
; APPLICANT: Cao, Yongwei
; APPLICANT: Wu, Wei
; APPLICANT: Boukharov, Andrey A.
; APPLICANT: Barbazuk, Brad
; APPLICANT: Li, Ping
; TITLE OF INVENTION: Rice Nucleic Acid Molecules and Other Molecules Associated With
; FILE OF INVENTION: Plants and Uses Thereof for Plant Improvement
; FILE REFERENCE: 38-21(53221)B
; CURRENT APPLICATION NUMBER: US/10/437,963
; CURRENT FILING DATE: 2003-05-14
; NUMBER OF SEQ ID NOS: 204966
; SEQ ID NO 8217
; LENGTH: 2185
; TYPE: DNA
; ORGANISM: Oryza sativa
; FEATURE:
; OTHER INFORMATION: Clone ID: PAT_MRT4530_14742C.1
US-10-437-963-8217

Alignment Scores:
Pred. No.: 564 Length: 2185
Score: 37.00 Matches: 7
Percent Similarity: 87.50% Conservative: 0
Best Local Similarity: 87.50% Mismatches: 1
Query Match: 90.24% Indels: 0

DB: 18 Gaps: 0
US-09-017-715A-2_COPY_120_127 (1-8) x US-10-437-963-8217 (1-2185)
Qy 1 GUGUUAAGInserGlyasp 8
Db 1208 GAGGAGCTCAGGAGGAGGTGAC 1231

Search completed: May 4, 2005, 16:39:46
Job time : 68.7405 secs

GenCore version 5.1.6
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OM protein - nucleic search, using frame_plus_p2n model

Run on: May 4, 2005, 09:07:55 ; Search time 18.5235 Seconds
(without alignments)
706.682 Million cell updates/sec

Title: US-09-017-715a-2_COPY_120_127

Perfect score: 41
Sequence: 1 EEAQSGSD 8

Scoring table:
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Ygapop 10.0, Ygapext 0.5
Fgapop 6.0, Fgapext 7.0
Delop 6.0, Delext 7.0

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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-LOOPTXT=0 -UNITS=bites -START=1 -END=-1 -MATRIX=blosum62 -TRANS=human40.cdi
-LIST=45 -DOCALLIGN=200 -THR SCORE=pct -THR MAX=100 -THR MIN=0 -ALIGN=15
-MODE=LOCAL -OUTFMT=p2n -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=200000000
-USER=US09017715_@CG2_1_116_@runat_04052005_100745_25632 -NCPU=6 -ICPU=3
-NO MMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG
-DEV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

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4: /cg2_6/ptodata/1/ina/6B.COMB.seq.*
5: /cg2_6/ptodata/1/ina/PCTUS.COMB.seq.*
6: /cg2_6/ptodata/1/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	41	100.0	550	US-08-705-771-1	Sequence 1, Appl1
2	41	100.0	550	US-09-417-540-1	Sequence 1, Appl1
3	41	100.0	702	US-09-949-016-1915	Sequence 1915, Ap
4	41	100.0	720	US-09-949-016-442	Sequence 442, App
5	36	87.8	1902	US-09-949-016-5550	Sequence 5550, Ap
6	36	87.8	4985	US-09-473-717-1	Sequence 1, Appl1
7	36	87.8	4985	US-09-949-016-152	Sequence 152, App
8	36	87.8	5515	US-09-398-193-98	Sequence 98, Appl
9	36	87.8	11214	US-09-949-016-17292	Sequence 17292, A
10	36	87.8	126200	US-09-949-016-11824	Sequence 11824, A
11	36	87.8	126200	US-09-949-016-13193	Sequence 13193, A
12	36	87.8	154605	US-09-949-016-11894	Sequence 11894, A

C 13	35	85.4	83708	4	US-09-949-016-17207	Sequence 17207, A
C 14	34	82.9	601	4	US-09-949-016-49739	Sequence 49739, A
C 15	34	82.9	628	3	US-09-328-111-367	Sequence 367, App
C 16	34	82.9	1653	4	US-09-830-111E-1	Sequence 1, Appl1
C 17	34	82.9	29717	4	US-09-949-016-16284	Sequence 16284, A
C 18	34	82.9	60376	4	US-09-949-016-12423	Sequence 12423, A
C 19	33	80.5	204	4	US-09-621-976-18715	Sequence 18715, A
C 20	33	80.5	210	4	US-09-621-976-18727	Sequence 18727, A
C 21	33	80.5	219	4	US-09-621-976-18782	Sequence 18782, A
C 22	33	80.5	229	4	US-09-621-976-18788	Sequence 18788, A
C 23	33	80.5	361	4	US-09-621-976-18728	Sequence 18728, A
C 24	33	80.5	404	4	US-09-621-976-18769	Sequence 18769, A
C 25	33	80.5	431	4	US-09-513-999C-13044	Sequence 13044, A
C 26	33	80.5	450	4	US-09-621-976-3657	Sequence 3657, App
C 27	33	80.5	480	4	US-09-270-767-6338	Sequence 6338, App
C 28	33	80.5	480	4	US-09-270-767-21620	Sequence 21620, A
C 29	33	80.5	487	4	US-09-621-976-18762	Sequence 18762, A
C 30	33	80.5	495	4	US-09-621-976-3659	Sequence 3659, App
C 31	33	80.5	601	4	US-09-949-016-26549	Sequence 26549, A
C 32	33	80.5	601	4	US-09-949-016-26550	Sequence 26550, A
C 33	33	80.5	601	4	US-09-949-016-27764	Sequence 27764, A
C 34	33	80.5	601	4	US-09-949-016-65608	Sequence 65608, A
C 35	33	80.5	601	4	US-09-949-016-65609	Sequence 65609, A
C 36	33	80.5	601	4	US-09-949-016-164928	Sequence 164928, A
C 37	33	80.5	601	4	US-09-949-016-202311	Sequence 202311, A
C 38	33	80.5	670	3	US-09-404-879A-156	Sequence 156, App
C 39	33	80.5	670	4	US-09-338-933-156	Sequence 156, App
C 40	33	80.5	670	4	US-09-215-681-156	Sequence 156, App
C 41	33	80.5	670	4	US-09-216-003A-156	Sequence 156, App
C 42	33	80.5	670	4	US-09-667-857-156	Sequence 156, App
C 43	33	80.5	978	4	US-09-248-796A-5309	Sequence 5309, App
C 44	33	80.5	1005	4	US-09-270-767-26942	Sequence 26942, A
C 45	33	80.5	1208	4	US-09-270-767-25543	Sequence 25543, A

ALIGNMENTS

RESULT 1
US-08-705-771-1
Sequence 1, Application US/08705771
Patent No. 6054289
GENERAL INFORMATION:
APPLICANT: Paul Moore, Reiner Gentz, Hongjin Ji.
TITLE OF INVENTION: Human Genes, Sequences and
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: CECCHI, STEWART & OLSTEIN
CITY: ROSELAND
STATE: NEW JERSEY
COUNTRY: USA
ZIP: 07068
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5 INCH DISKETTE
COMPUTER: IBM PS/2
OPERATING SYSTEM: MS-DOS
SOFTWARE: WORD PERFECT 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/705,771
FILING DATE: August 30, 1996
CLASSIFICATION: 536
ATTORNEY/AGENT INFORMATION:
NAME: MULLINS, J.G.
REGISTRATION NUMBER: 33,073
REFERENCE/DOCKET NUMBER: 325800-346 (PT196)
TELECOMMUNICATION INFORMATION:
TELEPHONE: 973-994-1700
TELEFAX: 973-994-1744
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:

LENGTH: 550 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-705-771-1

Alignment Scores:
Pred. No.: 9.29 Length: 550
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 3 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x US-08-705-771-1 (1-550)

Qy 1 GUGUUAAGInserG1yG1yasp 8
Db 369 GAGAGAGCCAGAGTGGGAGAC 392

RESULT 2
US-09-417-540-1
Sequence 1, Application US/09417540
Patent No. 6639052

GENERAL INFORMATION:
APPLICANT: Paul Moore, Reiner Gentz, Hongjin Ji,
Jian Ni and Jing-Shan Hu
TITLE OF INVENTION: Human Genes, Sequences and
Expression Products
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: CECCHIA, BYRNE, BAIN, GILFILLAN,
STREET: 6 BECKER FARM ROAD
CITY: ROSELAND
STATE: NEW JERSEY
COUNTRY: USA
ZIP: 07068

COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5 INCH DISKETTE
COMPUTER: IBM PS/2
OPERATING SYSTEM: MS-DOS
SOFTWARE: WORD PERFECT 5.1

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/417,540
FILING DATE: 14-OCT-1999
CLASSIFICATION: <Unknown>

PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/705,771
FILING DATE: August 30, 1996
ATTORNEY/AGENT INFORMATION:
NAME: MULLINS, J.G.
REGISTRATION NUMBER: 33,073
REFERENCE/DOCKET NUMBER: 325800-346 (PFI96)
TELECOMMUNICATION INFORMATION:
TELEPHONE: 973-994-1700
TELEFAX: 973-994-1744

INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 550 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-417-540-1

Alignment Scores:
Pred. No.: 9.29 Length: 550
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0

Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x US-09-417-540-1 (1-550)

Qy 1 GUGUUAAGInserG1yG1yasp 8
Db 369 GAGAGAGCCAGAGTGGGAGAC 392

RESULT 3
US-09-949-016-1915
Sequence 1915, Application US/09949016
Patent No. 6812339

GENERAL INFORMATION:
APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
FILE REFERENCE: CL001307
CURRENT APPLICATION NUMBER: US/09/949,016
PRIOR FILING DATE: 2000-04-14
PRIOR APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 60/231,498
PRIOR FILING DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 1915
LENGTH: 702
TYPE: DNA
ORGANISM: Human
US-09-949-016-1915

Alignment Scores:
Pred. No.: 12.1 Length: 702
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x US-09-949-016-1915 (1-702)

Qy 1 GUGUUAAGInserG1yG1yasp 8
Db 405 GAGAGAGCCAGAGTGGGAGAC 428

RESULT 4
US-09-949-016-442
Sequence 442, Application US/09949016
Patent No. 6812339

GENERAL INFORMATION:
APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
FILE REFERENCE: CL001307
CURRENT APPLICATION NUMBER: US/09/949,016
PRIOR FILING DATE: 2000-04-14
PRIOR APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 60/231,498
PRIOR FILING DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 442
LENGTH: 720
TYPE: DNA
ORGANISM: Human
US-09-949-016-442

Alignment Scores:

Pred. No.: 12, 4 Length: 720
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x US-09-949-016-442 (1-720)

QY 1 GluGluAlaGlnSerGlyIasp 8

DB 406 GAGGAGGCCAGAGTGGGGGAGAC 429

RESULT 5

US-09-949-016-5550/c
Sequence 5550, Application US/09949016
Patent No. 6812339
GENERAL INFORMATION:

APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

FILE REFERENCE: CL001307

CURRENT APPLICATION NUMBER: US/09/949, 016

CURRENT FILING DATE: 2000-04-14

PRIOR APPLICATION NUMBER: 60/241,755

PRIOR FILING DATE: 2000-10-20

PRIOR APPLICATION NUMBER: 60/237,768

PRIOR FILING DATE: 2000-10-03

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR FILING DATE: 2000-09-08

NUMBER OF SEQ ID NOS: 207012

SOFTWARE: FastSeq for Windows Version 4.0

SEQ ID NO 5550

LENGTH: 1902

TYPE: DNA

ORGANISM: Human

US-09-949-016-5550

Alignment Scores:

Pred. No.: 366 Length: 1902
Score: 36.00 Matches: 7
Percent Similarity: 87.50% Conservative: 0
Best Local Similarity: 87.50% Mismatches: 1
Query Match: 87.80% Indels: 0
DB: 4 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x US-09-949-016-5550 (1-1902)

QY 1 GluGluAlaGlnSerGlyIasp 8

DB 477 GAGGAGGCCAGCGCGGGGCGAC 454

RESULT 6

US-09-473-717-1/c
Sequence 1, Application US/09473717
Patent No. 6372475
GENERAL INFORMATION:

APPLICANT: Storm, Daniel R.
APPLICANT: Hacker, Beth
APPLICANT: Tomlinson, James E.

APPLICANT: COR Therapeutics, Inc.

TITLE OF INVENTION: University of Washington

TITLE OF INVENTION: CLONING AND CHARACTERIZATION OF A HUMAN ADENYLYL

FILE REFERENCE: 44481-5029-01-US

CURRENT APPLICATION NUMBER: US/09/473, 717

CURRENT FILING DATE: 1999-12-29

PRIOR APPLICATION NUMBER: PCT/US98/13541

PRIOR FILING DATE: 1998-07-01

PRIOR APPLICATION NUMBER: 60/098,559

PRIOR FILING DATE: 1997-07-01

PRIOR APPLICATION NUMBER: 08/886,440

PRIOR FILING DATE: 1997-07-01

NUMBER OF SEQ ID NOS: 3

SOFTWARE: PatentIn Ver. 2.0

SEQ ID NO 1

LENGTH: 4985

TYPE: DNA

ORGANISM: human type IX adenylyl cyclase

FEATURE:

NAME/KEY: CDS

LOCATION: (17)..(3898)

US-09-473-717-1

Alignment Scores:

Pred. No.: 1,04e+03 Length: 4985
Score: 36.00 Matches: 7
Percent Similarity: 87.50% Conservative: 0
Best Local Similarity: 87.50% Mismatches: 1
Query Match: 87.80% Indels: 0
DB: 3 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x US-09-473-717-1 (1-4985)

QY 1 GluGluAlaGlnSerGlyIasp 8

DB 475 GAGGAGGCCAGCGCGGGGCGAC 452

RESULT 7

US-09-949-016-152/c
Sequence 152, Application US/09949016
Patent No. 6812339
GENERAL INFORMATION:

APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

FILE REFERENCE: CL001307

CURRENT APPLICATION NUMBER: US/09/949, 016

CURRENT FILING DATE: 2000-04-14

PRIOR APPLICATION NUMBER: 60/241,755

PRIOR FILING DATE: 2000-10-20

PRIOR APPLICATION NUMBER: 60/237,768

PRIOR FILING DATE: 2000-10-03

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR FILING DATE: 2000-09-08

NUMBER OF SEQ ID NOS: 207012

SOFTWARE: FastSeq for Windows Version 4.0

SEQ ID NO 152

LENGTH: 4985

TYPE: DNA

ORGANISM: Human

US-09-949-016-152

Alignment Scores:

Pred. No.: 1,04e+03 Length: 4985
Score: 36.00 Matches: 7
Percent Similarity: 87.50% Conservative: 0
Best Local Similarity: 87.50% Mismatches: 1
Query Match: 87.80% Indels: 0
DB: 4 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x US-09-949-016-152 (1-4985)

QY 1 GluGluAlaGlnSerGlyIasp 8

DB 475 GAGGAGGCCAGCGCGGGGCGAC 452

RESULT 8

US-09-398-193-98/c
Sequence 98, Application US/09398193
Patent No. 6197581
GENERAL INFORMATION:

APPLICANT: Medical Research Council
TITLE OF INVENTION: Adenylylate cyclase and uses therefor
FILE REFERENCE: P24360-

```

; CURRENT APPLICATION NUMBER: US/09/398,193
; CURRENT FILING DATE: 1999-09-17
; NUMBER OF SEQ ID NOS: 104
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 98
; LENGTH: 5515
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (539)..(4600)
; US-09-398-193-98

Alignment Scores:
Pred. No.: 1,16e+03 Length: 5515
Score: 36.00 Matches: 7
Percent Similarity: 87.50% Conservative: 0
Best Local Similarity: 87.50% Mismatches: 1
Query Match: 87.80% Indels: 0
DB: 3 Gaps: 0

US-09-017-715a-2_copy_120_127 (1-8) x US-09-398-193-98 (1-5515)

QY 1 GUGUUAAGlnSerGlyGlyASP 8
DB 997 GAGGAAGCACAGCGCGGGGCGAC 974

RESULT 9
US-09-949-016-17292/c
; Sequence 17292, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 17292
; LENGTH: 112114
; TYPE: DNA
; ORGANISM: Human
; US-09-949-016-17292

Alignment Scores:
Pred. No.: 2.98e+04 Length: 112114
Score: 36.00 Matches: 7
Percent Similarity: 87.50% Conservative: 0
Best Local Similarity: 87.50% Mismatches: 1
Query Match: 87.80% Indels: 0
DB: 4 Gaps: 0

US-09-017-715a-2_COPY_120_127 (1-8) x US-09-949-016-17292 (1-112114)

QY 1 GUGUUAAGlnSerGlyGlyASP 8
DB 2477 GAGGAAGCACAGCGCGGGGCGAC 2454

RESULT 10
US-09-949-016-11824/c
; Sequence 11824, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
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; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 98
; LENGTH: 5515
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)...(126200)
; OTHER INFORMATION: n = A,T,C or G
; US-09-949-016-11824

Alignment Scores:
Pred. No.: 3.39e+04 Length: 126200
Score: 36.00 Matches: 7
Percent Similarity: 87.50% Conservative: 0
Best Local Similarity: 87.50% Mismatches: 1
Query Match: 87.80% Indels: 0
DB: 4 Gaps: 0

US-09-017-715a-2_COPY_120_127 (1-8) x US-09-949-016-11824 (1-126200)

QY 1 GUGUUAAGlnSerGlyGlyASP 8
DB 110246 GAGGAGCACAGTGGGGGGGAT 110223

RESULT 11
US-09-949-016-13193/c
; Sequence 13193, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13193
; LENGTH: 126200
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(126200)
; OTHER INFORMATION: n = A,T,C or G
; US-09-949-016-13193

Alignment Scores:
Pred. No.: 3.39e+04 Length: 126200
Score: 36.00 Matches: 7
Percent Similarity: 87.50% Conservative: 0
Best Local Similarity: 87.50% Mismatches: 1
Query Match: 87.80% Indels: 0
DB: 4 Gaps: 0
```


US-09-017-715A-2_COPY_120_127 (1-8) x US-09-949-016-13193 (1-126200)

QY 1 GluGluaLagInserGlyGlyAap 8

DB 110246 GAGGAGGACACAGCTGGGGGGAT 110223

RESULT 12

US-09-949-016-11894/C
; Sequence 11894, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CLO01307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 11894
; LENGTH: 154605
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-11894

Alignment Scores:

Pred. No.:	4.21e+04	Length:	154605
Score:	36.00	Matches:	7
Percent Similarity:	87.50%	Conservative:	0
Best Local Similarity:	87.50%	Mismatches:	1
Query Match:	87.80%	Indels:	0
DB:	4	Gaps:	0

US-09-017-715A-2_COPY_120_127 (1-8) x US-09-949-016-11894 (1-154605)

QY 1 GluGluaLagInserGlyGlyAap 8

DB 2475 GAGGAGGACACAGCTGGGGGGAT 2452

RESULT 13

US-09-949-016-17207/C
; Sequence 17207, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CLO01307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 17207
; LENGTH: 83708
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-17207

Alignment Scores:

Pred. No.:	3.46e+04	Length:	83708
Score:	35.00	Matches:	7

Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	85.37%	Indels:	0
DB:	4	Gaps:	0

US-09-017-715A-2_COPY_120_127 (1-8) x US-09-949-016-17207 (1-83708)

QY 1 GluGluaLagInserGlyGly 7

DB 2492 GAGGAGGACACAGCTGGGGGGAT 2472

RESULT 14

US-09-949-016-49739/C
; Sequence 49739, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CLO01307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 49739
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-49739

Alignment Scores:

Pred. No.:	268	Length:	601
Score:	34.00	Matches:	7
Percent Similarity:	87.50%	Conservative:	0
Best Local Similarity:	87.50%	Mismatches:	1
Query Match:	82.93%	Indels:	0
DB:	4	Gaps:	0

US-09-017-715A-2_COPY_120_127 (1-8) x US-09-949-016-49739 (1-601)

QY 1 GluGluaLagInserGlyGlyAap 8

DB 56 GAGGAGGACACAGCTGGGGGGAT 33

RESULT 15

US-09-328-111-367
; Sequence 367, Application US/09328111
; Patent No. 626233
; GENERAL INFORMATION:
; APPLICANT: Endege, Wilson O.
; APPLICANT: Steinmann, Kathleen E.
; APPLICANT: Ascle, Jon H.
; APPLICANT: Burgess, Christopher C.
; APPLICANT: Bushnell, Steven E.
; APPLICANT: Carroll III, Eddie
; APPLICANT: Catino, Theodore J.
; APPLICANT: Dertl, Adnan
; APPLICANT: Ford, Donna M.
; APPLICANT: Lewis, Marcia E.
; APPLICANT: Monahan, John E.
; APPLICANT: Schlegel, Robert
; TITLE OF INVENTION: NOVEL HUMAN GENES AND GENE EXPRESSION
; TITLE OF INVENTION: PRODUCTS
; FILE REFERENCE: CCD-257 (US)
; CURRENT APPLICATION NUMBER: US/09/328,111
; CURRENT FILING DATE: 1999-06-08
; EARLIER APPLICATION NUMBER: US 60/088,801

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; EARLIER FILING DATE: 1998-06-10
; NUMBER OF SEQ ID NOS: 850
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 367
; LENGTH: 628
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(628)
; OTHER INFORMATION: n = A,T,C or G
US-09-328-111-367

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Alignment Scores:
Pred. No.: 281          Length: 628
Score: 34.00          Matches: 6
Percent Similarity: 87.50%          Conservative: 1
Best Local Similarity: 75.00%          Mismatches: 1
Query Match: 82.93%          Indels: 0
DB: 3          Gaps: 0

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US-09-017-715A-2_COPY_120_127 (1-8) x US-09-328-111-367 (1-628)

```

Qy      1 GIUGIUAIGInSerGIYGIYasp 8
      ::||| ||| ||| ||| ||| ||| |||
Db      75 CAGGAAGCACACGACGAGGAGAGAC 98

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Search completed: May 4, 2005, 09:43:33
Job time : 63.5235 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - nucleic search, using frame_plus_p2n model

Run on: May 4, 2005, 10:17:21 ; Search time 1033.05 Seconds

(without alignments)
748,404 Million cell updates/sec

Title: US-09-017-715A-2

Perfect score: 610

Sequence: 1 MDVFKKGFSTAKKGVGAVG.....EGASKEKEVAEASQSGD 127

Scoring table:

BLOSUM62

Xgapop 10.0 , Xgapext 0.5

Ygapop 10.0 , Ygapext 0.5

Fgapop 6.0 , Fgapext 7.0

Delpop 6.0 , Delpext 7.0

Searched: 5642217 seqs, 3043843248 residues

Total number of hits satisfying chosen parameters: 11284434

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Command line parameters:

-MODEL=frame+g2n.model -DEV=x1h

-Q=/cgn2_1/USPRO.spool.h/US0901715/runat_04052005.100747.25719/app_query.fasta.1.661

-DB=Published Applications NA -QFMT=fastcap -SUFFIX=rnpb -MINMATCH=0.1

-LOOPCL=0 -LOOPEXT=0 -UNITS=bits -START=1 -END=1 -MATRIX=blonum62

-TRANS=human40.cdd -LIST=45 -DOCALIGN=200 -THR SCORE=pct -THR MAX=100

-THR MIN=0 -ALIGN=15 -MODE=LOCAL -OUTFMT=pct -NORM=ext -HEAPSIZE=500 -MINLEN=0

-MATELEN=200000000 -USER=US0901715 @CGN 1.1.684 @runat_04052005.100747.25719

-NCPU=6 -ICPU=3 -NO MMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSELOCK=100

-LONGLOG -DEV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5

-FGAPOP=6 -FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

Published Applications NA:*

1: /cgn2_6/ptodata/1/pubpna/US07_PUBCOMB.seq:*

2: /cgn2_6/ptodata/1/pubpna/PCT_NEM_PUB.seq:*

3: /cgn2_6/ptodata/1/pubpna/US06_NEM_PUB.seq:*

4: /cgn2_6/ptodata/1/pubpna/US06_PUBCOMB.seq:*

5: /cgn2_6/ptodata/1/pubpna/US07_NEM_PUB.seq:*

6: /cgn2_6/ptodata/1/pubpna/PCTUS_PUBCOMB.seq:*

7: /cgn2_6/ptodata/1/pubpna/US08_NEM_PUB.seq:*

8: /cgn2_6/ptodata/1/pubpna/US08_PUBCOMB.seq:*

9: /cgn2_6/ptodata/1/pubpna/US09A_PUBCOMB.seq:*

10: /cgn2_6/ptodata/1/pubpna/US09B_PUBCOMB.seq:*

11: /cgn2_6/ptodata/1/pubpna/US09C_PUBCOMB.seq:*

12: /cgn2_6/ptodata/1/pubpna/US09_NEM_PUB.seq:*

13: /cgn2_6/ptodata/1/pubpna/US10A_PUBCOMB.seq:*

14: /cgn2_6/ptodata/1/pubpna/US10B_PUBCOMB.seq:*

15: /cgn2_6/ptodata/1/pubpna/US10C_PUBCOMB.seq:*

16: /cgn2_6/ptodata/1/pubpna/US10D_PUBCOMB.seq:*

17: /cgn2_6/ptodata/1/pubpna/US10E_PUBCOMB.seq:*

18: /cgn2_6/ptodata/1/pubpna/US10F_PUBCOMB.seq:*

19: /cgn2_6/ptodata/1/pubpna/US10G_NEM_PUB.seq:*

20: /cgn2_6/ptodata/1/pubpna/US10H_NEM_PUB.seq:*

21: /cgn2_6/ptodata/1/pubpna/US11_NEM_PUB.seq:*

22: /cgn2_6/ptodata/1/pubpna/US60_PUBCOMB.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	610	100.0	550	9 US-09-954-531-613	Sequence 613, App
2	610	100.0	550	17 US-10-453-478-1	Sequence 1, Appl
3	610	100.0	550	19 US-10-843-641A-1680	Sequence 1680, Ap
4	602	98.7	796	4 US-09-925-286-171	Sequence 171, App
5	602	98.7	796	14 US-10-102-806-171	Sequence 171, App
6	595	97.5	384	19 US-10-826-157-5	Sequence 5, Appl
7	595	97.5	720	14 US-10-097-420-297	Sequence 297, App
8	592	97.0	720	17 US-10-282-174-469	Sequence 469, App
9	592	97.0	720	19 US-10-600-009-469	Sequence 469, App
10	538	88.2	479	10 US-09-918-995-2705	Sequence 2705, Ap
11	468.5	76.8	786	14 US-10-267-849-1	Sequence 1, Appl
12	328	53.8	210	18 US-10-204-337A-5	Sequence 1, Appl
13	316	51.8	1018	17 US-10-152-319A-1710	Sequence 1710, Ap
14	316	51.8	1018	19 US-10-486-706-260	Sequence 260, App
15	312.5	51.2	437	18 US-10-737-262-2	Sequence 2, Appl
16	307.5	50.4	423	14 US-10-077-584-1	Sequence 1, Appl
17	307.5	50.4	423	18 US-10-204-337A-3	Sequence 3, Appl
18	307.5	50.4	423	18 US-10-826-157-1	Sequence 1, Appl
19	307.5	50.4	423	19 US-10-737-262-1	Sequence 1, Appl
20	307.5	50.4	1105	15 US-10-223-978-10	Sequence 10, Appl
21	307.5	50.4	1466	15 US-10-101-510-362	Sequence 362, App
22	307.5	50.4	1543	18 US-10-721-693-14	Sequence 14, Appl
23	307.5	50.4	1543	18 US-10-852-997-14	Sequence 14, Appl
24	304.5	49.9	755	17 US-10-112-944-11	Sequence 11, Appl
25	304	49.8	441	9 US-09-960-352-12619	Sequence 12619, A
26	304	49.8	453	9 US-09-960-352-12629	Sequence 12629, Ap
27	303.5	49.8	1096	10 US-09-921-406C-35	Sequence 35, Appl
28	303.5	49.8	1096	18 US-10-721-693-23	Sequence 23, Appl
29	303.5	49.8	1096	18 US-10-852-997-23	Sequence 23, Appl
30	302	49.5	405	19 US-10-826-157-3	Sequence 3, Appl
31	253	41.5	249	18 US-10-204-337A-4	Sequence 4, Appl
32	243	39.8	424	9 US-09-960-352-1978	Sequence 1978, Ap
33	233	38.2	473	10 US-09-918-995-1832	Sequence 1832, Ap
34	205	33.6	456	9 US-09-918-995-26977	Sequence 26977, A
35	202	33.1	363	9 US-09-960-352-3369	Sequence 3369, Ap
36	201.5	33.0	666	17 US-10-240-425-388	Sequence 388, App
37	201.5	33.0	5666	17 US-10-282-174-72	Sequence 72, Appl
38	201.5	33.0	5666	17 US-10-600-009-72	Sequence 72, Appl
39	201.5	33.0	6012	17 US-10-282-174-483	Sequence 483, App
40	201.5	33.0	6012	17 US-10-600-009-483	Sequence 483, App
41	199	32.6	1332	16 US-10-029-386-25796	Sequence 25796, A
42	199	32.6	502	16 US-10-029-386-12096	Sequence 12096, A
43	197.5	32.4	5666	17 US-10-282-174-73	Sequence 73, Appl
44	197.5	32.4	5666	19 US-10-600-009-73	Sequence 73, Appl
45	195	32.0	521	16 US-10-029-386-9757	Sequence 9757, Ap

ALIGNMENTS

RESULT 1
US-09-954-531-613
Sequence 613, Application US/09954531
Patent No. US20020165180A1
GENERAL INFORMATION:
APPLICANT: Weaver, Zoe
TITLE OF INVENTION: Process for Identifying Anti-Cancer Therapeutic Agents Using Canc
FILE REFERENCE: 689290-77
CURRENT FILING DATE: 2002-05-02
PRIOR APPLICATION NUMBER: US/09/954, 531
PRIOR FILING DATE: 2000-09-18
PRIOR APPLICATION NUMBER: US/60/233, 133
PRIOR FILING DATE: 2000-09-20
PRIOR APPLICATION NUMBER: US/60/234, 009
PRIOR FILING DATE: 2000-09-20
PRIOR APPLICATION NUMBER: US/60/234, 034
PRIOR FILING DATE: 2000-09-20
PRIOR APPLICATION NUMBER: US/60/234, 509
PRIOR FILING DATE: 2000-09-22
PRIOR APPLICATION NUMBER: US/60/234, 567

PRIOR FILING DATE: 2000-09-22
NUMBER OF SEQ ID NOS: 1392
SOFTWARE: PatentIn version 3.0
SEQ ID NO 613
LENGTH: 550
TYPE: DNA
ORGANISM: Homo sapiens
US-09-954-531-613

Alignment Scores:
Pred. No.: 2,15e-60 Length: 550
Score: 610.00 Matches: 127
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0

US-09-017-715a-2 (1-127) x US-09-954-531-613 (1-550)

QY 1 MetAspValPheLysGlyPheSerIleAlaLysGlyValValGluValGlu 20
DB 12 ATGGATGTTTCAAGAGGGCTTCTCCATCGCCAGAGGGCGTGGTGGCGGGA 71

QY 21 LysThrLysGlnGlyValThrGluAlaGluLysThrLysGlnGlyValMetCyrVal 40
DB 72 AAGACCAAGCAGGGGTGACGAGACGCTGAGAACCAAGAGGGGTGATGTATGTG 131

QY 41 G1ValAlaLysThrLysGluAsnValValGlnSerValThrSerValAlaGluLysThrLys 60
DB 132 GGAGGCCAAGACCAAGAGATGTGTACAGACGCTGACCTCGGCGGAGAACCAAG 191

QY 61 GluGlnAlaAsnAlaValSerLysAlaValAlaSerSerValAsnThrValAlaThrLys 80
DB 192 GAGCAGGCCAAGCCGTGACAGGCTGTGTGAGAGGCTCAACCTGTGGCCACCAAG 251

QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValValArgLysGluAspLeu 100
DB 252 ACCGTGAGAGAGCGGAGAACATCGCGTCACCTCGGGGTGTGCGCAAGAGGACTTG 311

QY 101 ArgProSerAlaProGlnGlnGlnGluLysAlaSerLysGluLysGlnGluValAlaGlu 120
DB 312 AGGCCATCTGCCCCCAACAGAGGGGTGAGGCATCCAAAGAAAGAGAAAGTGGCAGAG 371

QY 121 GluAlaGlnSerGlyGlyAsp 127
DB 372 GAGGCCCAAGAGTGGGGGAGAC 392

RESULT 2
US-10-453-478-1
Sequence 1, Application US/10453478
Publication No. US20030208043A1
GENERAL INFORMATION:
APPLICANT: Paul Moore, Reiner Gentz, Hongjin Ji,
Jian Ni and Jing-Shan Hu
TITLE OF INVENTION: Human Genes, Sequences and
Expression Products
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: CARRELLA, BYRNE, BAIN, GILFILLAN,
CECCHI, STEWART & OLSTEIN
STREET: 6 BECKER FARM ROAD
CITY: ROSELAND
STATE: NEW JERSEY
COUNTRY: USA
ZIP: 07068
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5 INCH DISKETTE
COMPUTER: IBM PS/2
OPERATING SYSTEM: MS-DOS
SOFTWARE: WORD PERFECT 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/10/453,478
FILING DATE: 04-Jun-2003

CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/08/705,771
FILING DATE: August 30, 1996
ATTORNEY/AGENT INFORMATION:
NAME: MULLINS, J.G.
REGISTRATION NUMBER: 33,073
REFERENCE/DOCKET NUMBER: 325800-346 (PFI96)
TELECOMMUNICATION INFORMATION:
TELEPHONE: 973-994-1700
TELEFAX: 973-994-1744
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 550 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-10-453-478-1

Alignment Scores:
Pred. No.: 2,15e-60 Length: 550
Score: 610.00 Matches: 127
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 17 Gaps: 0

US-09-017-715a-2 (1-127) x US-10-453-478-1 (1-550)

QY 1 MetAspValPheLysGlyPheSerIleAlaLysGlyValValGluValGlu 20
DB 12 ATGGATGTTTCAAGAGGGCTTCTCCATCGCCAGAGGGCGTGGTGGCGGGA 71

QY 21 LysThrLysGlnGlyValThrGluAlaGluLysThrLysGlnGlyValMetCyrVal 40
DB 72 AAGACCAAGCAGGGGTGACGAGACGCTGAGAACCAAGAGGGGTGATGTATGTG 131

QY 41 G1ValAlaLysThrLysGluAsnValValGlnSerValThrSerValAlaGluLysThrLys 60
DB 132 GGAGGCCAAGACCAAGAGATGTGTACAGACGCTGACCTCGGCGGAGAACCAAG 191

QY 61 GluGlnAlaAsnAlaValSerLysAlaValAlaSerSerValAsnThrValAlaThrLys 80
DB 192 GAGCAGGCCAAGCCGTGACAGGCTGTGTGAGAGGCTCAACCTGTGGCCACCAAG 251

QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValValArgLysGluAspLeu 100
DB 252 ACCGTGAGAGAGCGGAGAACATCGCGTCACCTCGGGGTGTGCGCAAGAGGACTTG 311

QY 101 ArgProSerAlaProGlnGlnGlnGluLysAlaSerLysGluLysGlnGluValAlaGlu 120
DB 312 AGGCCATCTGCCCCCAACAGAGGGGTGAGGCATCCAAAGAAAGAGAAAGTGGCAGAG 371

QY 121 GluAlaGlnSerGlyGlyAsp 127
DB 372 GAGGCCCAAGAGTGGGGGAGAC 392

RESULT 3
US-10-843-641A-1680
Sequence 1680, Application US/10843641A
Publication No. US20050064454A1
GENERAL INFORMATION:
APPLICANT: Avalon Pharmaceuticals, Inc.
TITLE OF INVENTION: Cancer Gene Determination and Therapeutic Screening Using
File Reference: 689290-189
FILE REFERENCE: Signature Gene Sets
CURRENT APPLICATION NUMBER: US/10/843,641A
CURRENT FILING DATE: 2004-05-12
PRIOR APPLICATION NUMBER: US/09/873,367
PRIOR FILING DATE: 2001-06-05
PRIOR APPLICATION NUMBER: US/09/954,531

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CURRENT FILING DATE: 2001-08-10
PRIOR APPLICATION NUMBER: PCT/US00/05881
PRIOR FILING DATE: 2000-03-08
PRIOR APPLICATION NUMBER: 60/124,270
PRIOR FILING DATE: 1999-03-12
NUMBER OF SEQ ID NOS: 846
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 171
LENGTH: 796
TYPE: DNA
ORGANISM: Homo sapiens
US-09-925-298-171

Alignment Scores:
Pred. No.: 2,87e-59 Length: 796
Score: 602.00 Matches: 125
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 98.43% Mismatches: 0
Query Match: 98.63% Indels: 0
DB: Gaps: 0

US-09-017-715A-2 (1-127) x US-09-925-298-171 (1-796)

QY 1 MetaSpValPhylLyLyGlyPheSerTLeAlaYsLySGlyValAlaGlyAlaGlu 20
|||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||
Db 109 ATGAGATGYTTTCAGAGAGGCGCTTCTCCATCCGCCAAGGAGGCGCTGCGTG3AA 168

QY 21 LysThrLySGlnGlyValThrGluAlaAlaGluThrLySGlnGlyValMetYrVal 40
|||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||
Db 169 AGACCAAGCAGAGGGGTATCGAGACAGCTAGAGAGCAAGAGAGGGGTCTATATGTG 228

QY 41 GYAlaLysThrLySGluAsnValAlaGlnSerValThrSerValAlaGluYrThrLys 60
|||:::|||||:::|||||:::|||||:::|||||:::|||||
Db 229 GAGGCCAAGACCAAGAGATGTGTATACAGACGCTGACCTCATGTGCGCGAAGACCAAG 288

QY 61 GluGlnAlaAsnAlaValSerLysAlaValSerSerValAsnThrValAlaThrLys 80
|||:::|||||:::|||||:::|||||:::|||||:::|||||
Db 289 GAGCAGAGCCCAAGCCGTGAGGAGCGTGTGTGAGAGCGTCGACACTGTGGCCACCAAG 348

QY 81 ThrValGluGluAlaGluAsnLeAlaValThrSerGlyValAlaTyrLySGluAspLeu 100
|||:::|||||:::|||||:::|||||:::|||||:::|||||
Db 349 ACCGTGAGAGAGCGCGAAGACATCGCGTCACTCCGGGGTGTGTCCCAAGAGACTTG 408

QY 101 ArgProSerAlaProGlnGlnGluGlyGluAlaSerLySGluYrSGluGluValAlaGlu 120
|||:::|||||:::|||||:::|||||:::|||||:::|||||
Db 409 AGGCCATCTGCCCCCAACAGAGAGGTATGAGCATCCAAAGAGAAAGAGAAATGGCAGAG 468

QY 121 GluAlaGlnSerGlyGlyYrAsp 127
|||:::|||||
Db 469 GAGGCCCAAGCTGGCGGAGAC 489

RESULT 5
US-10-102-806-171
Sequence 171, Application US/10102806
Publication No. US20030054421A1
GENERAL INFORMATION:
APPLICANT: Rosen et al.
TITLE OF INVENTION: Nucleic Acids, Proteins and Antibodies
FILE REFERENCE: PA103P1C1
CURRENT APPLICATION NUMBER: US/10/102,806
CURRENT FILING DATE: 2002-03-22
PRIOR APPLICATION NUMBER: 09/925,298
PRIOR FILING DATE: 2001-08-10
PRIOR APPLICATION NUMBER: PCT/US00/05881
PRIOR FILING DATE: 2000-03-08
PRIOR APPLICATION NUMBER: 60/124,270
PRIOR FILING DATE: 1999-03-12
NUMBER OF SEQ ID NOS: 846
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 171
LENGTH: 796
TYPE: DNA
ORGANISM: Homo sapiens

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US-10-102-806-171

Alignment Scores:

Pred. No.: 2,87e-59 Length: 796
Score: 602.00 Matches: 125
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 98.43% Mismatches: 0
Query Match: 98.69% Indels: 0
DB: 14 Gaps: 0

US-09-017-715A-2 (1-127) x US-10-102-806-171 (1-796)

QY 1 MetAspValPheLysGlyPheSerIleAlaLysGlyValValGlyValAlaGlu 20
DB 109 ATGATGCTTTCAAGAGGCGCTTCTCCATCGCCAGAGAGGGGTGGTGGCGGTGGA 168
QY 21 LysThrLysGlnGlyValThrGluAlaAglLysThrLysGlnGlyValMetCysVal 40
DB 169 AAGACCAAGCAGGGGGTGCAGAGCAGCTGAGAAAGACCAAGAGGGGTCTCATGTATGTG 228
QY 41 G1A1A1LysThrLysGlnAsnValValGlnSerValThrSerValAlaGluLysThrLys 60
DB 229 GGAGCCAAAGACCAAGAGATGTGTACAGACGTGACCTCAGTGGCCGAGAAACCAAG 288
QY 61 G1G1A1A1AsnAlaValSerLysAlaValLysSerValAsnThrValAlaThrLys 80
DB 289 GAGCAGGCCAAGCGCGTGCAGAGCTGTGTGAGCAGCGTCAACACTGTGGCCACCAAG 348
QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValValArgLysGluAspLeu 100
DB 349 ACCGTGAGAGAGCGGAGAACATCGCGCTCACCTCCGGGGTGGTGGCAAGAGGACTTG 408
QY 101 ArgProSerAlaProGlnGlnGlnGlyGluLysGlnLysGluGluValAlaGlu 120
DB 409 AGGCCATCTGCCCCCAACAGAGGGGTGAGGCATCCAAAGAAAGAGAGTGGCAGG 468
QY 121 G1uA1aGlnSerGlyGlyAsp 127
DB 469 GAGGCCCAAGAGTGGGGGAGAC 489

RESULT 6

US-10-826-157-5
; Sequence 5, Application US/10826157
; Publication No. US20050064548A1
; GENERAL INFORMATION:
; APPLICANT: Lindquist, Susan L.
; APPLICANT: Outeiro, Tiago
; TITLE OF INVENTION: YEAST ECTOPOICALLY EXPRESSING ABNORMALLY
; FILE REFERENCE: 17481-003001
; CURRENT APPLICATION NUMBER: US/10/826,157
; CURRENT FILING DATE: 2004-04-16
; PRIOR APPLICATION NUMBER: US 60/472,317
; PRIOR FILING DATE: 2003-05-20
; PRIOR APPLICATION NUMBER: US 60/463,284
; NUMBER OF SEQ ID NOS: 8
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 5
; LENGTH: 384
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-826-157-5

Alignment Scores:

Pred. No.: 6,95e-59 Length: 384
Score: 595.00 Matches: 124
Percent Similarity: 99.21% Conservative: 2
Best Local Similarity: 97.64% Mismatches: 1
Query Match: 97.54% Indels: 0
DB: 19 Gaps: 0

US-09-017-715A-2 (1-127) x US-10-826-157-5 (1-384)

QY 1 MetAspValPheLysGlyPheSerIleAlaLysGlyValValGlyValAlaGlu 20
DB 1 ATGATGCTTTCAAGAGGCGCTTCTCCATCGCCAGAGAGGGGTGGTGGCGGTGGA 60
QY 21 LysThrLysGlnGlyValThrGluAlaAglLysThrLysGlnGlyValMetCysVal 40
DB 61 AAGACCAAGCAGGGGGTGCAGAGCAGCTGAGAAAGACCAAGAGGGGTCTCATGTATGTG 120
QY 41 G1A1A1LysThrLysGlnAsnValValGlnSerValThrSerValAlaGluLysThrLys 60
DB 121 GGAGCCAAAGACCAAGAGATGTGTACAGACGTGACCTCAGTGGCCGAGAAACCAAG 180
QY 61 G1G1A1A1AsnAlaValSerLysAlaValLysSerValAsnThrValAlaThrLys 80
DB 181 GAGCAGGCCAAGCGCGTGCAGAGCTGTGTGAGCAGCGTCAACACTGTGGCCACCAAG 240
QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValValArgLysGluAspLeu 100
DB 241 ACCGTGAGAGAGCGGAGAACATCGCGGTCACTCCGGGGTGGTGGCAAGAGGACTTG 300
QY 101 ArgProSerAlaProGlnGlnGlnGlyGluLysGlnLysGluGluValAlaGlu 120
DB 301 AGGCCATCTGCCCCCAACAGAGGGGTGAGGCATCCAAAGAAAGAGAGTGGCAGAG 360
QY 121 G1uA1aGlnSerGlyGlyAsp 127
DB 361 GAGGCCCAAGAGTGGGGGAGAC 381

RESULT 7

US-10-097-340-297
; Sequence 297, Application US/10097340
; Publication No. US20030087250A1
; GENERAL INFORMATION:
; APPLICANT: John MONAHAN
; APPLICANT: Manjula GANNAVARAPU
; APPLICANT: Sebastian HOERSCH
; APPLICANT: Shubhangi KAMATKAR
; APPLICANT: Steve G. KOVATS
; APPLICANT: Rachel E. MEYERS
; APPLICANT: Michael MORRISSEY
; APPLICANT: Peter OLANDT
; APPLICANT: Ami SEN
; APPLICANT: Peter VEIBY
; APPLICANT: Gordon B. MILLS
; APPLICANT: Robert C. BAST, Jr.
; APPLICANT: Karen LU
; APPLICANT: Rosemarie SCHMANDT
; APPLICANT: Xumei ZHAO
; TITLE OF INVENTION: Nucleic Acid Molecules and Proteins For The Identification,
; FILE REFERENCE: Assessment, Prevention, and Therapy of Ovarian Cancer
; CURRENT APPLICATION NUMBER: US/10/097,340
; CURRENT FILING DATE: 2002-03-14
; PRIOR APPLICATION NUMBER: 60/276,025
; PRIOR FILING DATE: 2001-03-14
; PRIOR APPLICATION NUMBER: 60/325,149
; PRIOR FILING DATE: 2001-09-26
; PRIOR APPLICATION NUMBER: 60/276,026
; PRIOR FILING DATE: 2001-03-14
; PRIOR APPLICATION NUMBER: 60/324,967
; PRIOR FILING DATE: 2001/09/26
; PRIOR APPLICATION NUMBER: 60/311,732
; PRIOR FILING DATE: 2001-08-10
; PRIOR APPLICATION NUMBER: 60/325,102
; PRIOR FILING DATE: 2001-09-26
; PRIOR APPLICATION NUMBER: 60/323,580
; NUMBER OF SEQ ID NOS: 363
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 297
; LENGTH: 720

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; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-097-340-297

Alignment Scores:
Pred. No.:      1,59e-58          Length:      720
Score:          595.00           Matches:     124
Percent Similarity: 99.21%       Conservative: 2
Best Local Similarity: 97.64%    Mismatches: 1
Query Match:     97.54%         Indels:     0
DB:              14             Gaps:       0

US-09-017-715A-2 (1-127) x US-10-097-340-297 (1-720)

QY      1 MetAdPValPhelyelysGlyPheSerIleAlaLysGlyValValGlyAlaValGlu 20
Db      49 ATGATGTCTTCACAGAAAGGGCTTCTCATCGCCAAAGGGCGTGCGTGCAGAA 108
        |||
QY      21 LyeThrLyvGlnGlyValThrGluAlaAlaGluLysThrLysGluGlyValMetCylrVal 40
Db      109 AAGACCACCAAGCAGGGGTGACGGAACGACTGAAACAACCAAGAGGGCGCTCATGTGTG 168
        |||
QY      41 GlyAlaLysThyLysGluLysValValGlnSerValThrservalAlaGluLysThryls 60
Db      169 GGAGGCCAACAGCACAGAGAATGTGTRACAGAGCTTAACCTCACTGCGCCGAAAGACCAAG 228
        |||
QY      61 GluGlnAlaAsnAlaValserLysAlaValAlaSerServalAsnthrValAlatrlys 80
Db      229 GAGCAGGCGCAACGCGGTGAGCAGCGCTGTGTGACGAGCTCAACACTGTGGCCACCAAG 288
        |||
QY      81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValValArgLysGluAspleu 100
Db      289 ACCGTGGAGGAGGAGCGAGAAATCGGGTCACCTCGGGGGTGTGTGGCAAAGGAGACTTG 348
        |||
QY      101 ArgProSerAlaProGlngInglnglGlyGluAlaSerLysGluLysGluGluValAlaglu 120
Db      349 AGCCCATCTGCCCCCCCACAGAGAGGTGTGCATCCAAAGAAAGAGAGTAGTGACAG 408
        |||
QY      121 GluAlaGlnSerGlyGlyASP 127
Db      409 GAGGCCACAGATGGGGGAGAGAC 429
        |||

RESULT 8
US-10-282-174-469
; Sequence 469, Application US/10282174
; Publication No. US20030224380A1
GENERAL INFORMATION:
APPLICANT: Becker, Kenneth David
APPLICANT: Velicelebi, Gonul
APPLICANT: Eliot, Kathryn J.
APPLICANT: Wang, Xin
APPLICANT: Tanzi, Rudolph E.
APPLICANT: Bertam, Lars
APPLICANT: Saunders, Aleister J.
APPLICANT: Mullin, Kristina M.
APPLICANT: Sampson, Andrew Johnson
TITLE OF INVENTION: Blacker, Deborah Lynne
TITLE OF INVENTION: GENES AND POLYMORPHISMS ON CHROMOSOME 10
TITLE OF INVENTION: ASSOCIATED WITH ALZHEIMER'S DISEASE AND OTHER
FILE REFERENCE: 37481-3308
CURRENT APPLICATION NUMBER: US/10/282,174
PRIOR FILING DATE: 2002-10-25
PRIOR APPLICATION NUMBER: US 60/339,525
PRIOR FILING DATE: 2001-10-25
PRIOR APPLICATION NUMBER: US 60/338,010
PRIOR FILING DATE: 2001-11-08
PRIOR APPLICATION NUMBER: US 60/336,929
PRIOR FILING DATE: 2001-11-08
PRIOR APPLICATION NUMBER: US 60/338,363
PRIOR FILING DATE: 2001-11-09
PRIOR APPLICATION NUMBER: US 60/337,052
PRIOR FILING DATE: 2001-12-04

```

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P R I O R   A P P L I C A T I O N   N U M B E R :   U S   6 0 / 3 6 8 , 9 1 9
P R I O R   F I L I N G   D A T E :   2 0 0 2 - 0 3 - 2 8
N U M B E R   O F   S E Q   I D   N O S :   5 6 4
S O F T W A R E :   F A S T S E Q   f o r   W i n d o w s   V e r s i o n   4 . 0
S E Q   I D   N O   4 6 9
L E N G T H :   7 2 0
T Y P E :   D N A
O R G A N I S M :   H o m o   s a p i e n s
F E A T U R E :
N A M E / K E Y :   a l l e l e
L O C A T I O N :   3 0 , 5 7 , 8 5 , 2 4 3 , 2 5 0 , 3 7 7 , 5 1 2 , 5 3 1 , 5 5 5 , 5 6 1 , 6 7 2
O T H E R   I N F O R M A T I O N :   N   i s   a n y
U S - 1 0 - 2 8 2 - 1 7 4 - 4 6 9

A l i g n m e n t   S c o r e s :
P r e d .   N o . :           3 , 5 6 - 5 8           L e n g t h :           7 2 0
S c o r e :                   5 9 2 . 0 0           M a t c h e s :           1 2 4
P e r c e n t   S i m i l a r i t y :   9 7 . 6 4 %   C o n s e r v a t i v e :   0
B e s t   L o c a l   S i m i l a r i t y :   9 7 . 6 4 %   M i s m a t c h e s :   3
Q u e r y   M a t c h :           9 7 . 0 5 %         I n d e l s :           0
D B :                           1 7               G a p s :           0

U S - 0 9 - 0 1 7 - 7 1 5 A - 2   ( 1 - 1 2 7 )   x   U S - 1 0 - 2 8 2 - 1 7 4 - 4 6 9   ( 1 - 7 2 0 )

Q Y      1   M e t a b P a l P h e l y s G l y P h e S e r I e a l a l y L y s G l y V a l a l g l y a l a V a l g l u   2 0
D b      4 9   A T G A T G T T T C C A G A A G G G C T T C C A T C C G C C A A G N A G G G G T G T G G T G C G T G G A   1 0 8
Q Y      2 1   L y s T h r L y s G l n G l V a l T h r G l u n l a l a g l u l y s T h r L y s G l n g l y V a l M e t T y r V a l   4 0
D b      1 0 9   A A G A C C A A G C A G G G G G T A C G A A C A C T G T G A A G A C C A A G A G G G G G C A T G T A T G T G   1 6 8
Q Y      4 1   G l y a l a l y S t h r L y s G l u a s n l a l a V a l G l n s e r V a l T h r S e r V a l a l a g l u l y s T h r L y s   6 0
D b      1 6 9   G G A G C C A A G A C C A A G A G A A T T T T G T A C A G A C G T G A C C T C A G T G C C G A A G A C C A A G   2 2 8
Q Y      6 1   G l u G l n a l a a s n l a l a V a l S e r L y s a l a V a l a l S e r S e r V a l a s n T h r V a l a t h r L y s   8 0
D b      2 2 9   G A G C A G G C C A A C G C N G T A G A G C T G T G T G A C A G C G T C A A C A C T G T G C C A C C A A G   2 8 8
Q Y      8 1   T h r V a l g l u g l u a l a g l u a s n l l e a l a V a l T h r S e r G l y V a l a r g l y s G l u a p l e u   1 0 0
D b      2 8 9   A C C G G A G A G A G G C G A A A C T C C G G T C A C T C C G G G G T G T G C G A A G A G A G A C T T G   3 4 8
Q Y      1 0 1   A r g P r o S e r a l a P r o G l n G l n g l u g l u a l S e r L y s G l u l y s G l u l y s G l u V a l a l a g l u   1 2 0
D b      3 4 9   A G C C A T T G C C C C C C C A C A G A G G G T N G C A T C C A A A G A A A G A G A A G T G C A G   4 0 8
Q Y      1 2 1   G l u a l a g l n s e r G l y l a s p   1 2 7
D b      4 0 9   G A G G C C A G A G T G G G G G A G A C   4 2 9

R E S U L T   9
U S - 1 0 - 6 0 0 - 0 0 9 - 4 6 9
S e q u e n c e   4 6 9 ,   A p p l i c a t i o n   U S / 1 0 6 0 0 0 0 9
P u b l i c a t i o n   N o .   U S 2 0 0 5 0 0 9 3 0 3 1 a 1
G E N E R A L   I N F O R M A T I O N :
A P P L I C A N T :   B e c k e r ,   K e n n e t h   D a v i d
A P P L I C A N T :   V e l i c e l e b i ,   G o n u l
A P P L I C A N T :   E l l i o t ,   K a t h r y n   J .
A P P L I C A N T :   W a n g ,   X i n
A P P L I C A N T :   T a n z i ,   R u d o l p h   E .
A P P L I C A N T :   B e r t r a m ,   L a r s
A P P L I C A N T :   S a u n d e r s ,   A l e i s t e r   J .
A P P L I C A N T :   M u l l i n ,   K r i s t i n a   M .
A P P L I C A N T :   S a m p s o n ,   A n d r e w   J o h n s o n
A P P L I C A N T :   B l a c k e r ,   D e b o r a h   L y n n e
T I T L E   O F   I N V E N T I O N :   G E N E S   A N D   P O L Y M O R P H I S M S   O N   C H R O M O S O M E   1 0
T I T L E   O F   I N V E N T I O N :   A S S O C I A T E D   W I T H   A L Z H E I M E R ' S   D I S E A S E   A N D   O T H E R
F I L E   R E F E R E N C E :   3 7 4 8 1 - 3 3 0 8 B
C U R R E N T   A P P L I C A T I O N   N U M B E R :   U S / 1 0 / 6 0 0 . 0 0 9
C U R R E N T   F I L I N G   D A T E :   2 0 0 3 - 0 6 - 1 8

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; PRIOR APPLICATION NUMBER: US 60/339,525
; PRIOR FILING DATE: 2001-10-25
; PRIOR APPLICATION NUMBER: US 60/338,010
; PRIOR FILING DATE: 2001-11-08
; PRIOR APPLICATION NUMBER: US 60/336,929
; PRIOR FILING DATE: 2001-11-08
; PRIOR APPLICATION NUMBER: US 60/338,363
; PRIOR FILING DATE: 2001-11-09
; PRIOR APPLICATION NUMBER: US 60/337,052
; PRIOR FILING DATE: 2001-12-04
; PRIOR APPLICATION NUMBER: US 60/368,919
; PRIOR FILING DATE: 2002-03-28
; PRIOR APPLICATION NUMBER: US 10/282,174
; PRIOR FILING DATE: 2002-10-25
; NUMBER OF SEQ ID NOS: 564
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 469
; LENGTH: 720
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: allele
; LOCATION: 30,57,85,243,250,377,512,531,555,561,672
; OTHER INFORMATION: N is any
; US-10-600-009-469

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Alignment Scores:
Pred. No.: 3.5e-58 Length: 720
Score: 592.00 Matches: 124
Percent Similarity: 97.64% Conservative: 0
Best Local Similarity: 97.64% Mismatches: 3
Query Match: 97.05% Indels: 0
Gaps: 19

```

US-09-017-715A-2 (1-127) x US-10-600-009-469 (1-720)

```

QY 1 Metaspvalphelyslysglypheserilealyseglyvalvalglvalaaglu 20
DB 49 ATGAGTATGTTTCAAGAGGCGCTTCTCATCGCCAGNAGGCGGTGGTGGCGGAGAA 108
QY 21 LysThrLysGlnGlyValThrGluAlaGluLysThrLysGlnGlyValMetLysVal 40
DB 109 AAGACCAAGCAGGGGTGACGAGACAGCTGAGAACAGCAGAGGGGTGATGATG 168
QY 41 GAlaLysThrLysGlnGluAsnValAlaGlnSerValThrSerValAlaGluLysThrLys 60
DB 169 GAGCCAGACCAAGAGAGATGTTGACAGAGCTGACCTCGAGCCAGAACCAAG 228
QY 61 GlnGlnAlaAsnAlaValSerLysAlaValAlaSerSerValAsnThrValAlaThrLys 80
DB 229 GAGCAGGCCAAGCGNGTGAGCNAAGGCTGTGTGAGCAGCGTCAACACTGTGCCCAAG 288
QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValAlaArgLysGluAspLeu 100
DB 289 ACCGTGAGAGAGCGGAGAAATCGGCGTCACTCGGAGGTGGTGGCAAGAGAGCTTG 348
QY 101 AcgProSerAlaProGlnGlnGluGluAlaSerLysGluLysGlnGluValAlaGlu 120
DB 349 AGGCACTCTGCCCCCAAGAGAGGCTGCGGCTCCAAAGAGAAAGAGAGTGGCAGG 408
QY 121 GluAlaGlnSerGlyLysAsp 127
DB 409 GAGGCCCAAGAGTGGGGAAC 429

```

RESULT 10

```

US-09-918-995-2705
; Sequence 2705, Application US/09918995
; Publication No. US20030073623A1
; GENERAL INFORMATION:
; APPLICANT: HySeq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
; FILE REFERENCE: 20411-756

```

```

; CURRENT APPLICATION NUMBER: US/09/918,995
; CURRENT FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: US/09/235,076
; PRIOR FILING DATE: 1999-01-20
; NUMBER OF SEQ ID NOS: 38054
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 2705
; LENGTH: 479
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(479)
; OTHER INFORMATION: n = A,T,C or G
; US-09-918-995-2705

```

```

Alignment Scores:
Pred. No.: 3.03e-52 Length: 479
Score: 538.00 Matches: 112
Percent Similarity: 99.14% Conservative: 3
Best Local Similarity: 96.55% Mismatches: 1
Query Match: 88.20% Indels: 0
Gaps: 10

```

US-09-017-715A-2 (1-127) x US-09-918-995-2705 (1-479)

```

QY 12 LysLysGlyValAlaGluAlaValAlaGluLysThrLysGlnGlyValThrGluAlaGlu 31
DB 47 CAGAGGGCGGTGAGNGGGTGGCGGTGAGAAAGACCAAGCAGGGGTGACGGAAGCAGCTGAG 106
QY 32 LysThrLysGlnGlyValMetLysValAlaLysThrLysGluAsnValAlaGlnSer 51
DB 107 AAGACCAAGAGGGGTGATGATGTGGAGCAAGCAAGAGAAATGTTGTACAGAGC 166
QY 52 ValThrSerValAlaGluLysThrLysGlnGlnAlaAsnAlaValSerLysAlaValAla 71
DB 167 GTGACCTCAGTGGCCGAGAGACCAAGAGCAGCGCCAGCGCTGAGAGGCTGTGGTG 226
QY 72 SerSerValAsnThrValAlaThrLysThrValGluGluAlaGluAsnIleAlaValThr 91
DB 227 AGCAGCGTCAACACTGTGGCCCAAGACCGTGGAGAGAGGGCGGAGAACACTGGCGTCAAC 286
QY 92 SerGlyValAlaArgLysGluAspLeuArgProSerAlaProGlnGlnGluGluAla 111
DB 287 TCCGGGGTGGTGGCAGAGAGGACTTGAGGCCATCTGCCCCCAAGAGAGGTGAGCA 346
QY 112 SerLysGluLysGlnGluValAlaGluGlnGlnGlnGlnGlnGlnGlnGlnGlnGln 127
DB 347 TCCAAAGAGAAAGAGAGAGTGGCAGAGAGGCCCAAGAGTGGGGAGAC 394

```

RESULT 11

```

US-10-267-849-1
; Sequence 1, Application US/10267849
; Publication No. US20030087824A1
; GENERAL INFORMATION:
; APPLICANT: Ji, Hongjun
; APPLICANT: Rosen, Craig A.
; TITLE OF INVENTION: Breast Cancer Specific Gene 2
; FILE REFERENCE: 1488.0810001
; CURRENT APPLICATION NUMBER: US/10/267,849
; CURRENT FILING DATE: 2002-10-10
; PRIOR APPLICATION NUMBER: US/08/673,284
; PRIOR FILING DATE: 1996-06-28
; PRIOR APPLICATION NUMBER: US 60/000,602
; PRIOR FILING DATE: 1995-06-30
; NUMBER OF SEQ ID NOS: 45
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1
; LENGTH: 786
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-10-267-849-1

```


Alignment Scores:

Pred. No.: 5.09e-44 Length: 786
Score: 468.50 Matches: 111
Percent Similarity: 86.15% Conservative: 1
Best Local Similarity: 85.38% Mismatches: 15
Query Match: 76.80% Indels: 4
DB: 14 Gaps: 1

US-09-017-715A-2 (1-127) x US-10-267-849-1 (1-786)

Qy 1 MetaspVAlpHeLySgLyPheSerIleAlaLySgLyValValGlyAlaValGlu 20
Db 95 ATGATGTCTTCAAGAGGGCTTCTCCATGCCAAGAGGGCGTGGCGGTGGAA 154
Qy 21 LySgThLySgLnGlyValThrGluAlaGluLySgThLySgLnGlyValMetTyVal 40
Db 155 AAGACCAAGAGGGGGTGGAGGAGCGTGAAGAGCAAGAGGGGGTCACTATATGTG 214
Qy 41 GlyAlaLySgThLySgLnGluAen---Val-ValGlnSerValThSerValAlaGluLySgTh 59
Db 215 GAGGCCAAGACCAAGAGGAATGTGTATGTCAGAGGCTCACTGCGCCGAGAAC 274
Qy 59 rLySgLnGluAlaAenAlaValSerLySgAlaValSerSerValAenThrValAlaTh 79
Db 275 CAAGGAGCAGGCGCAACCGCTGAGCAAGCGTGTGTGAGCAGCGTCAACACTKTGGCCAC 334
Qy 79 rLySgThValGluGluAlaGluAenIleAlaValThSerGlyValValArgLySgLnG 99
Db 335 CAAGACCGTGGAGGAGGCGGAGAACATCGCGTCACTCCGGGTGTGTCGCAAGAGGA 394
Qy 99 pLeuArgProSerAlaProGlnGlnGluGlyGluAlaSerLySgLn-LySgLnGluVala 119
Db 395 YTKKAGGCCATTY-TKCCCCAAGCAGAGGGTGGAGGCTCAARAGARARAKMGSAAAGWG 453
Qy 119 laGluGluAlaGlnSerGlyGlyAap 127
Db 454 CMRAKKRGKMSGAGAGTGGGGAGAC 479

RESULT 12

US-10-204-337A-5
; Sequence 5, Application US/10204337A
; Publication No. US20040128706A1
; GENERAL INFORMATION:
; APPLICANT: Maelib, Eliezer
; TITLE OF INVENTION: Method for screening for Anti-Amyloidogenic Properties and Method
; FILE REFERENCE: 6627-PC9014
; CURRENT APPLICATION NUMBER: US/10/204,337A
; PRIOR FILING DATE: 2002-08-16
; PRIOR APPLICATION NUMBER: US 60/183,571
; PRIOR FILING DATE: 2000-02-18
; PRIOR APPLICATION NUMBER: PCT/US00/07216
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 5
; LENGTH: 210
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-204-337A-5

Alignment Scores:

Pred. No.: 1.02e-28 Length: 210
Score: 328.00 Matches: 68
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 97.14% Mismatches: 0
Query Match: 53.77% Indels: 0
DB: 18 Gaps: 0

US-09-017-715A-2 (1-127) x US-10-204-337A-5 (1-210)

Qy 1 MetaspVAlpHeLySgLyPheSerIleAlaLySgLyValValGlyAlaValGlu 20
Db 28 ATGATGTCTTCAAGAGGGCTTCTCCATGCCAAGAGGGCGTGGCGGTGGAA 154

Db 1 ATGATGTCTTCAAGAGGGCTTCTCCATGCCAAGAGGGCGTGGCGGTGGAA 60
Qy 21 LySgThLySgLnGlyValThrGluAlaGluLySgThLySgLnGlyValMetTyVal 40
Db 61 AAGACCAAGAGGGGGTGGAGGAGCGTGAAGAGCAAGAGGGGGTCACTATATGTG 120
Qy 41 GlyAlaLySgThLySgLnGluAenValValGlnSerValThSerValAlaGluLySgTh 60
Db 121 GAGGCCAAGACCAAGAGGAATGTGTATGTCAGAGGCTCACTGCGCCGAGAACCAAG 180
Qy 61 GluGlnAlaAenAlaValSerLySgAlaVal 70
Db 181 GAGCAGGCCAAGCGGTGAGCAGAGCGTGTG 210

RESULT 13

US-10-152-319A-1710
; Sequence 1710, Application US/10152319A
; Publication No. US20040072160A1
; GENERAL INFORMATION:
; APPLICANT: Mendrick, Donna
; APPLICANT: Porter, Mark
; APPLICANT: Johnson, Kory
; APPLICANT: Hyspe, Brandon
; APPLICANT: Castle, Arthur
; APPLICANT: Blaschoff, Michael
; TITLE OF INVENTION: Molecular Toxicology Modeling
; FILE REFERENCE: 44921-5089-US
; CURRENT APPLICATION NUMBER: US/10/152,319A
; PRIOR FILING DATE: 2002-05-22
; PRIOR APPLICATION NUMBER: US 60/292,335
; PRIOR FILING DATE: 2001-05-22
; PRIOR APPLICATION NUMBER: US 60/297,523
; PRIOR FILING DATE: 2001-06-13
; PRIOR APPLICATION NUMBER: US 60/298,925
; PRIOR FILING DATE: 2001-06-19
; PRIOR APPLICATION NUMBER: US 60/303,810
; PRIOR FILING DATE: 2001-07-10
; PRIOR APPLICATION NUMBER: US 60/303,807
; PRIOR FILING DATE: 2001-07-10
; PRIOR APPLICATION NUMBER: US 60/303,808
; PRIOR FILING DATE: 2001-07-10
; PRIOR APPLICATION NUMBER: US 60/315,047
; PRIOR FILING DATE: 2001-08-28
; PRIOR APPLICATION NUMBER: US 60/324,928
; PRIOR FILING DATE: 2001-09-27
; PRIOR APPLICATION NUMBER: US 60/330,867
; PRIOR FILING DATE: 2001-11-01
; PRIOR APPLICATION NUMBER: US 60/330,462
; PRIOR FILING DATE: 2001-10-22
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 2221
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1710
; LENGTH: 1018
; TYPE: DNA
; ORGANISM: Rattus norvegicus
; FEATURE:
; OTHER INFORMATION: Genbank Accession No. NM_019169
US-10-152-319A-1710

Alignment Scores:

Pred. No.: 1.91e-26 Length: 1018
Score: 316.00 Matches: 70
Percent Similarity: 68.60% Conservative: 13
Best Local Similarity: 57.85% Mismatches: 32
Query Match: 51.80% Indels: 6
DB: 17 Gaps: 1

US-09-017-715A-2 (1-127) x US-10-152-319A-1710 (1-1018)

Qy 1 MetaspVAlpHeLySgLyPheSerIleAlaLySgLyValValGlyAlaValGlu 20
Db 28 ATGATGTCTTCAAGAGGGCTTCTCCATGCCAAGAGGGCGTGGCGGTGGAA 154

QY 21 LysThrLysGlnGlyValThrGluAlaGluLysThrLysGlnGlyValMetTyrVal 40
DB 88 AAAACCAAGCAGGAGGCTGGCAAGCAGCTGGGAAGCAAAAGAGGCGCTCTATGTGA 147
QY 41 G1yAlaLysThrLysGlnGluSerValValGlnSerValThrSerValAlaGluLysThrLys 60
DB 148 GGTTCACAAACCTAAGAGGAGGAGTCTGTTCAATGAGACAAAGTGGCTGAGAAAGCAAA 207
QY 61 G1uGlnAlaAsnAlaValSerLysAlaValValSerSerValAsnThrValAlaThrLys 80
DB 208 GAGCAAGTACCAAAATGTTGAGAGGAGGAGTGTGATGACAGCACTCCCTCAGAAAG 267
QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValValArgLysGluAsnLys 100
DB 268 ACAGTGGAGGAGGAGCTGGGAACATTTGCTGCCACATGTTTGTCAAGAAAGACCAAGATG 327
QY 101 ArgProSerAlaProGlnGlnGluGluAlaSerLysGluLysGluGluValAlaGlu 120
DB 328 -----GCAAGGCTGAAGAAAGGATACCAAGAGGGAATCTCTGGA 369
QY 121 G1u 121
DB 370 GAC 372

RESULT 14

US-10-486-706-260
Sequence 260, Application US/10486706
Publication No. US20050071088A1
GENERAL INFORMATION:
APPLICANT: LANDFIELD, PHILIP W.
APPLICANT: BLALOCK, ERIC M.
APPLICANT: CHEN, KUEY-CHU
APPLICANT: FOSTER, THOMAS C.
TITLE OF INVENTION: GENE EXPRESSION PROFILE BIOMARKERS AND THERAPEUTIC TARGETS FOR
TITLE OF INVENTION: BRAIN AGING AND AGE-RELATED COGNITIVE IMPAIRMENT
FILE REFERENCE: 50229-426
CURRENT APPLICATION NUMBER: US/10/486,706
CURRENT FILING DATE: 2004-02-13
PRIOR APPLICATION NUMBER: PCT/US02/25607
PRIOR FILING DATE: 2002-08-13
PRIOR APPLICATION NUMBER: US 60/311,343
PRIOR FILING DATE: 2001-08-13
NUMBER OF SEQ ID NOS: 461
SOFTWARE: PatentIn version 3.2
SEQ ID NO: 260
LENGTH: 1018
TYPE: DNA
ORGANISM: Rattus norvegicus
US-10-486-706-260

Alignment Scores:

Pred. No.: 1,91e-26 Length: 1018
Score: 316.00 Matches: 70
Percent Similarity: 68.60% Conservative: 13
Best Local Similarity: 57.85% Mismatches: 32
Query Match: 51.80% Indels: 6
DB: 19 Gaps: 1

US-09-017-715A-2 (1-127) x US-10-486-706-260 (1-1018)

QY 1 MetAspValPheLysLysGlyPheSerIleAlaLysGlyValValGluValGlu 20
DB 28 ATGATGTGTTTCAAGAAAGACTTCAAGAGCCAGAGGAGTGTGCTGCTGAG 87
QY 21 LysThrLysGlnGlyValThrGluAlaGluLysThrLysGlnGlyValMetTyrVal 40
DB 88 AAAACCAAGCAGGAGGCTGGCAAGCAGCTGGGAAGCAAAAGAGGCGCTCTATGTGA 147
QY 41 G1yAlaLysThrLysGlnGluSerValValGlnSerValThrSerValAlaGluLysThrLys 60
DB 148 GGTTCACAAACCTAAGAGGAGGAGTCTGTTCAATGAGACAAAGTGGCTGAGAAAGCAAA 207

QY 61 G1uGlnAlaAsnAlaValSerLysAlaValValSerSerValAsnThrValAlaThrLys 80
DB 208 GAGCAAGTACCAAAATGTTGAGAGGAGGAGTGTGATGACAGCACTCCCTCAGAAAG 267
QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValValArgLysGluAsnLys 100
DB 268 ACAGTGGAGGAGGAGCTGGGAACATTTGCTGCCACATGTTTGTCAAGAAAGACCAAGATG 327
QY 101 ArgProSerAlaProGlnGlnGluGluAlaSerLysGluLysGluGluValAlaGlu 120
DB 328 -----GCAAGGCTGAAGAAAGGATACCAAGAGGGAATCTCTGGA 369
QY 121 G1u 121
DB 370 GAC 372

RESULT 15

US-10-737-262-2
Sequence 2, Application US/10737262
Publication No. US20040197315A1
GENERAL INFORMATION:
APPLICANT: Stefanie, Leonidas
APPLICANT: Greene, Lloyd A.
TITLE OF INVENTION: Dopaminergic Cell lines Stably Expressing A53T Alpha-Synuclein
TITLE OF INVENTION: and Methods of Using Same
FILE REFERENCE: 5199-26
CURRENT APPLICATION NUMBER: US/10/737,262
CURRENT FILING DATE: 2003-12-15
NUMBER OF SEQ ID NOS: 2
SOFTWARE: PatentIn version 3.2
SEQ ID NO: 2
LENGTH: 437
TYPE: DNA
ORGANISM: Homo sapiens
US-10-737-262-2

Alignment Scores:

Pred. No.: 1.58e-26 Length: 437
Score: 312.50 Matches: 69
Percent Similarity: 71.82% Conservative: 10
Best Local Similarity: 62.73% Mismatches: 28
Query Match: 51.23% Indels: 3
DB: 18 Gaps: 1

US-09-017-715A-2 (1-127) x US-10-737-262-2 (1-437)

QY 1 MetAspValPheLysLysGlyPheSerIleAlaLysGlyValValGluValGlu 20
DB 15 ATGATGTGTTTCAAGAAAGACTTCAAGAGCCAGAGGAGTGTGCTGCTGAG 74
QY 21 LysThrLysGlnGlyValThrGluAlaGluLysThrLysGlnGlyValMetTyrVal 40
DB 75 AAAACCAAGCAGGAGGCTGGCAAGCAGCTGGGAAGCAAAAGAGGCGTCTCTATGTGA 134
QY 41 G1yAlaLysThrLysGlnGluSerValValGlnSerValThrSerValAlaGluLysThrLys 60
DB 135 GGTTCACAAACCTAAGAGGAGGAGTGTGATGACAAAGTGGCTGAGAAAGCAAA 194
QY 61 G1uGlnAlaAsnAlaValSerLysAlaValValSerSerValAsnThrValAlaThrLys 80
DB 195 GAGCAAGTACCAAAATGTTGAGAGGAGGAGTGTGATGACAGCACTCCCTCAGAAAG 254
QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValValArgLysGluAsnLys 100
DB 255 ACAGTGGAGGAGGAGCTGGGAACATTTGCTGCCACATGTTTGTCAAGAAAGACCAAGATG 314
QY 101 -----ArgProSerAlaProGlnGln 107
DB 315 GCAAGAAATGAAGAGAGGCCCAACAGGA 344

Search completed: May 4, 2005, 16:39:31
Job time: 1042.71 secs

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GenCore version 5.1.6
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OM protein - nucleic search, using frame_plus_p2n model

Run on: May 4, 2005, 09:07:56 ; Search time 5778.07 Seconds
(without alignments) 836.639 Million cell updates/sec

Title: US-09-017-715A-2

Perfect score: 610
Sequence: 1 MDVFKKGFSJAKKGVGAVE.....EGEASKEKEVAEBAQSGSD 127

Scoring table:
BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68473088

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Command line parameters:

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-Q=/cgn2_1/USPTO.epool_h/US09017715/runat_04052005_100744_25619/app_query.fasta_1.661
-DB=EST -QPM=fastap -SUFFIX=rest -MINMATCH=0.1 -LOOPCL=0 -LOPEXT=0
-UNITS=bits -START=1 -END=1 -MATRIX=blomsum62 -TRANS=human40.cdi -LIST=45
-OUTFMT=pcio -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=200000000
-USER=US09017715_@CGN_1_1_5334_@runat_04052005_100744_25619 -NCPU=6 -ICPU=3
-NO MMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG
-DEV TIMEOUT=120 -MAIN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

EST: *
1: gb_est1.*
2: gb_est2.*
3: gb_hic.*
4: gb_est3.*
5: gb_est4.*
6: gb_est5.*
7: gb_est6.*
8: gb_gse1.*
9: gb_gse2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match Length	ID	Description
1	602	98.7	555	7 CV028548 7090 Full1
2	602	98.7	578	5 BP212912 BP212912
3	602	98.7	582	5 BP197662 BP197662
4	602	98.7	584	5 BP201686 BP201686
5	602	98.7	653	4 BP175131 BP175131
6	602	98.7	751	4 BP183596 BP183596
7	602	98.7	781	5 BP089395 BP089395
8	602	98.7	809	5 BP921124 BP921124
9	602	98.7	855	5 BP0882072 BP0882072

10	602	98.7	884	5 BP039430 BP039430
11	602	98.7	891	5 BP0221776 BP0221776
12	602	98.7	903	4 BP286466 BP286466
13	595	97.5	558	4 BP154891 BP154891
14	595	97.5	568	4 BP0708703 BP0708703
15	595	97.5	617	4 BP0707764 BP0707764
16	595	97.5	659	7 CN410061 CN410061
17	595	97.5	755	4 BP1597796 BP1597796
18	595	97.5	824	4 BP1600882 BP1600882
19	591	96.9	765	4 BP1603159 BP1603159
20	591	96.9	768	4 BP1603171 BP1603171
21	589	96.6	937	5 BP157619 BP157619
22	589	96.6	1009	5 BP179779 BP179779
23	576	94.4	641	1 AL712443 AL712443
24	575	94.3	555	5 BX474500 BX474500
25	567	93.0	623	5 BP381244 BP381244
26	566.5	92.9	799	4 BP1488930 BP1488930
27	564.5	92.5	949	5 BP066800 BP066800
28	557	91.3	706	4 BG328738 BG328738
29	555	91.0	583	5 BP200612 BP200612
30	548	89.8	583	5 BP346497 BP346497
31	542	88.9	756	5 BP0901053 BP0901053
32	541	88.7	489	5 BP201709 BP201709
33	526	86.2	462	5 BX474511 BX474511
34	524	85.9	467	6 CB107161 CB107161
35	523	85.7	511	2 AM659211 AM659211
36	523	85.7	625	7 CR452251 CR452251
37	523	85.7	625	7 CR454933 CR454933
38	523	85.7	646	7 CR551748 CR551748
39	523	85.7	670	7 CO879176 CO879176
40	523	85.7	689	7 CR454179 CR454179
41	523	85.7	700	7 CK770094 CK770094
42	523	85.7	801	7 CO873848 CO873848
43	522.5	85.7	721	7 CN157077 CN157077
44	522.5	85.7	721	7 CN159030 CN159030
45	519	85.1	799	7 CO877016 CO877016

ALIGNMENTS

RESULT 1
CV028548
LOCUS
DEFINITION 7090 Full Length cDNA from the Mammalian Gene Collection Homo sapiens CDNA 5' similar to BC014098, mRNA sequence.
ACCESSION CV028548
VERSION CV028548.1 GI:51486632
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM
REFERENCE
AUTHORS
Rual J.F., Hirozane-Kishikawa T., Hao T., Bertin N., Li S., Clingingsmith T.R., Hartley J.L., Dosescu D., Cheo D., Moore T., Simmons B., Sequerra R., Bosak S., Boudette-Stamm L., Le Pench C., Vandenhaute J., Cusick M.E., Alcala J.S., Hill D.E. and Vidal M. Human ORFeome Version 1.1: a Platform for Reverse Proteomics Genome Res. (2004) in press
Contact: Vidal M

TITLE JOURNAL
COMMENT
Marc Vidal Laboratory
Dana Farber Cancer Institute
1 Jimmy Fund Way Smith 858, BOSTON, MA 02115, USA
Tel: 617 632 5180
Fax: 617 632 5739
Email: Marc.Vidal@dfci.harvard.edu
ORF Sequence Tag (OST) of Gateway Entry construct. Each cloned ORF results from a PCR reaction using an MGC full-length cDNA as template DNA and ORF specific primers
PCR Primers
FORWARD: ATGATGTCTTCAAGAAGGCTTCTC
BACKWARD: TAGTCTCCCACTCTGG

Insert Length: 555 Std Error: 48.00
 Seq: 11017 row: 05 column: B
 Sled primer: ACTGCCCTCCTTACACGTCGTACTCGGAAAC
 High quality sequence start: 97
 High quality sequence stop: 554
 POLYA=No.

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 /tissue_type="mixed"
 /clone_lib="Full Length cDNA from the Mammalian Gene Collection"
 /note="Vector: mixed; The ORFs were PCR amplified from the MGC (Mammalian Gene Collection) as of April 2004 and cloned by recombinational Gateway cloning into pDONR223 Donor vector. Reference : MGC (Mammalian Gene Collection) Program Team, Generation and Initial Analysis of more than 15,000 Full-length Human and Mouse cDNA Sequences. PNAS, 2002, 99(126), 16899-16903"

ORIGIN

Alignment Scores:
 Pred. No.: 1,39e-58 Length: 555
 Score: 602.00 Matches: 125
 Percent Similarity: 100.00% Conservative: 2
 Best Local Similarity: 98.43% Mismatches: 0
 Query Match: 98.69% Indels: 0
 DB: 7 Gaps: 0

US-09-017-715a-2 (1-127) x CVO28548 (1-555)

QY 1 MetAepValPheLyseLyseLyseSerIleAlaLyseLyseGlyValAlaGlyAlaValAlaGlu 20
 DB 1 ATGGAGTCTTCAGAAAAGGGCTTCTCCATCGCCAGAGAGGGCGTGGTGGGTGGGAGAA 60
 QY 21 LysThrLysGlnGlyValThrGluAlaAlaGluLysThrLysGlnGlyValMetCyrVal 40
 DB 61 AAGACCAAGACAGGGGGGTGACGAGACAGCTGAGAACACCAAGAGGGGCTCATGTATG 120
 QY 41 GlyAlaLysThrLysGlnLysValAlaGlnSerValThrSerValAlaGluLysThrLys 60
 DB 121 GGAGCCAGACCAAGAGATGTTGTACAGAGCGTCACTCACTGCCAGAAAGACCAAG 180
 QY 61 GluGlnAlaAsnAlaValSerLysAlaValAlaSerSerValAsnThrValAlaThrLys 80
 DB 181 GAGCAGGCCAACGCCGTGAGCGAGCGTGTGTAGACAGCGCTCAACACTGTGGCCACCAAG 240
 QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValAlaArgLysGluAspLeu 100
 DB 241 ACCGTGAGAGAGCGGAGAACATCGCGGTCACTCCGGGGTGTGGCCAAAGAGAGACTTG 300
 QY 101 ArgProSerAlaProGlnGlnGlnGlyGluAlaSerLysGluLysGlnGluValAlaGlu 120
 DB 301 AGGCCATCTGCCCGCCCAACAGAGGGGTGAGGCATCCAAAGAAAGAGAGAGTGGCAGAG 360
 QY 121 GluAlaGlnSerGlyLysAsp 127
 DB 361 GAGGCCACAGATGGGGAGAC 381

RESULT 2
 BP12912 578 bp mRNA linear EST 15-SEP-2004
 LOCUS BP12912 Sugano cDNA library, cerebrum Homo sapiens cDNA clone
 DEFINITION CBR05118, mRNA sequence.
 ACCESSION BP12912
 VERSION BP12912.1 GI:52085803
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Cranialata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 578)
 AUTHORS Suzuki,Y., Yamashita,R., Shiota,M., Sakakibara,Y., Chiba,J.,
 Mizushima-Sugano,J., Nakai,K. and Sugano,S.
 TITLE Sequence comparison of human and mouse genes reveals a homologous
 block structure in the promoter regions
 JOURNAL Genome Res. 14 (9), 1711-1718 (2004)
 COMMENT Contact: Yutaka Suzuki
 Department of Virology
 Institute of Medical Science, University of Tokyo
 4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
 Email: ysuzuki@ims.u-tokyo.ac.jp.

FEATURES
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 1..578
 Location/Qualifiers
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ORIGIN

Alignment Scores:
 Pred. No.: 1,47e-58 Length: 578
 Score: 602.00 Matches: 125
 Percent Similarity: 100.00% Conservative: 2
 Best Local Similarity: 98.43% Mismatches: 0
 Query Match: 98.69% Indels: 0
 DB: 5 Gaps: 0

US-09-017-715a-2 (1-127) x BP12912 (1-578)

QY 1 MetAepValPheLyseLyseLyseSerIleAlaLyseLyseGlyValAlaGlyAlaValAlaGlu 20
 DB 125 ATGGAGTCTTCAGAAAAGGGCTTCTCCATCGCCAGAGAGGGCGTGGTGGGTGGGAGAA 184
 QY 21 LysThrLysGlnGlyValThrGluAlaAlaGluLysThrLysGlnGlyValMetCyrVal 40
 DB 185 AAGACCAAGACAGGGGGGTGACGAGACAGCTGAGAACACCAAGAGGGGCTCATGTATG 244
 QY 41 GlyAlaLysThrLysGlnLysValAlaGlnSerValThrSerValAlaGluLysThrLys 60
 DB 245 GGAGCCAGACCAAGAGATGTTGTACAGAGCGTCACTCACTGCCAGAAAGACCAAG 304
 QY 61 GluGlnAlaAsnAlaValSerLysAlaValAlaSerSerValAsnThrValAlaThrLys 80
 DB 305 GAGCAGGCCAACGCCGTGAGCGAGCGTGTGTAGACAGGTCAACACTGTGCCACCAAG 364
 QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValAlaArgLysGluAspLeu 100
 DB 365 ACCGTGAGAGAGCGGAGAACATCGCGGTCACTCCGGGGTGTGGCCAAAGAGAGACTTG 424
 QY 101 ArgProSerAlaProGlnGlnGlnGlyGluAlaSerLysGluLysGlnGluValAlaGlu 120
 DB 425 AGGCCATCTGCCCGCCCAACAGAGGGGTGAGGCATCCAAAGAAAGAGAGAGTGGCAGAG 484
 QY 121 GluAlaGlnSerGlyLysAsp 127
 DB 485 GAGGCCACAGATGGGGAGAC 505

RESULT 3
 BP197662 582 bp mRNA linear EST 14-SEP-2004
 LOCUS BP197662 Sugano cDNA library, adrenal gland Homo sapiens cDNA clone
 DEFINITION ADG06551, mRNA sequence.
 ACCESSION BP197662
 VERSION BP197662.1 GI:52043849
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Cranialata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 582)
 REFERENCE Suzuki,Y., Yamashita,R., Shiota,M., Sakakibara,Y., Chiba,J.,

TITLE Mizushima-Sugano, J., Nakai, K. and Sugano, S.
Sequence comparison of human and mouse genes reveals a homologous
block structure in the promoter regions
JOURNAL Genome Res. 14 (9), 1711-1718 (2004)
COMMENT Contact: Yutaka Suzuki
Department of Virology
Institute of Medical Science, University of Tokyo
4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
Email: yuzuki@ims.u-tokyo.ac.jp.

FEATURES
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1. 582
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ORIGIN

Alignment Scores:
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Score: 602.00 Matches: 125
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 98.43% Mismatches: 0
Query Match: 98.69% Indels: 0
DB: 5 Gaps: 0

US-09-017-715a-2 (1-127) x BP197662 (1-582)

QY 1 MetApyAlPheLySgLyPheSerLeAlaLySgLyValAlGlyAlaValAlGlu 20
DB 85 ATGATGCTCTTCAAGAGGCGCTTCTCATGCCAAGAGGCGGTGGGTGGAA 144
QY 21 LyThrLySgInGlyValThrGluAlaAlaGluLyThrLySgInGlyValMetTyVal 40
DB 145 AAGACCAAGAGGGGGTGACGAGAGCTGAGAAAGCAAGAGGGGGTCATGATGTG 204
QY 41 GYAlaLyThrLySgInGluAenValValGInSerValThrSerValAlaGluLyThrLyS 60
DB 205 GAGGCCAAGACCAAGAGAAATGTTGTACAGCGTGAAGCTGAGCCGAGAAACCAAG 264
QY 61 GluGlnAlaAenAlaValSerLySgAlaValSerSerValAenThrValAlaThrLyS 80
DB 265 GAGCAGGCCAAGCGCGTGAAGAGCGTGTGTGACGCGTCAACACTGTGCCACCAAG 324
QY 81 ThrValGluGluAlaGluAenAlaValThrSerGlyValValAArgLySgInGluApyLeu 100
DB 325 ACCGTGAGAGGAGCGGAGAACATCGCGTCACTCCGGGGTGTGCGCAAGAGGACTTG 384
QY 101 ArgProSerAlaProGInGInGInGlyGluAlaSerLySgInGluGluValAlaGlu 120
DB 385 AGGCACTCTGCCCCCAAGAGAGGTGAGGCAATCCAAAGAAAGAGAGTGGCAGAG 444
QY 121 GluAlaGInSerGlyGlyAap 127
DB 445 GAGGCCAGAGTGGGGAGAGC 465

RESULT 4
BP201686 584 bp mRNA linear EST 14-SEP-2004
LOCUS BP201686 Sugano CDNA library, amygdala Homo sapiens CDNA clone
DEFINITION AMR06165, mRNA sequence.
ACCESSION BP201686
VERSION BP201686.1 GI:52051909
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 584)
AUTHORS Suzuki, Y., Yamashita, R., Shirota, M., Sakakibara, Y., Chiba, J.,
Mizushima-Sugano, J., Nakai, K. and Sugano, S.
TITLE Sequence comparison of human and mouse genes reveals a homologous

JOURNAL block structure in the promoter regions
COMMENT Genome Res. 14 (9), 1711-1718 (2004)
Contact: Yutaka Suzuki
Department of Virology
Institute of Medical Science, University of Tokyo
4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
Email: yuzuki@ims.u-tokyo.ac.jp.

FEATURES
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1. 584
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/mol_type="mRNA"
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ORIGIN

Alignment Scores:
Pred. No.: 1,496-58 Length: 584
Score: 602.00 Matches: 125
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 98.43% Mismatches: 0
Query Match: 98.69% Indels: 0
DB: 5 Gaps: 0

US-09-017-715a-2 (1-127) x BP201686 (1-584)

QY 1 MetApyAlPheLySgLyPheSerLeAlaLySgLyValAlGlyAlaValAlGlu 20
DB 125 ATGATGCTCTTCAAGAGGCGCTTCTCATGCCAAGAGGCGGTGGGTGGAA 184
QY 21 LyThrLySgInGlyValThrGluAlaAlaGluLyThrLySgInGlyValMetTyVal 40
DB 185 AAGACCAAGAGGGGGTGACGAGAGCTGAGAAAGCAAGAGGGGGTCATGATGTG 244
QY 41 GYAlaLyThrLySgInGluAenValValGInSerValThrSerValAlaGluLyThrLyS 60
DB 245 GAGGCCAAGACCAAGAGAAATGTTGTACAGCGTGAAGCTGAGCCGAGAAACCAAG 304
QY 61 GluGlnAlaAenAlaValSerLySgAlaValSerSerValAenThrValAlaThrLyS 80
DB 305 GAGCAGGCCAAGCGCGTGAAGAGCGTGTGTGACGCGTCAACACTGTGCCACCAAG 364
QY 81 ThrValGluGluAlaGluAenAlaValThrSerGlyValValAArgLySgInGluApyLeu 100
DB 365 ACCGTGAGAGGAGCGGAGAACATCGCGTCACTCCGGGGTGTGCGCAAGAGGACTTG 424
QY 101 ArgProSerAlaProGInGInGInGlyGluAlaSerLySgInGluGluValAlaGlu 120
DB 425 AGGCACTCTGCCCCCAAGAGAGGTGAGGCAATCCAAAGAAAGAGAGTGGCAGAG 484
QY 121 GluAlaGInSerGlyGlyAap 127
DB 485 GAGGCCAGAGTGGGGAGAGC 505

RESULT 5
B1757131 653 bp mRNA linear EST 25-SEP-2001
LOCUS B1757131
DEFINITION 603030894F1 NIH_MGC_114 Homo sapiens CDNA clone IMAGE:520162 5',
mRNA sequence.
ACCESSION B1757131
VERSION B1757131.1 GI:15748709
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE NIH-MGC http://mgc.nci.nih.gov/
AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
TITLE Unpublished (1999)
JOURNAL Contact: Robert Strauberg, Ph.D.
COMMENT Email: cga@rs-remail.nih.gov

Tissue Procurement: Life Technologies, Inc.
 cDNA Library Preparation: Life Technologies, Inc.
 DNA Sequencing by: The I.M.A.G.E. Consortium (LLNL)
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
 Plate: L1M11503 row: 1 column: 23
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 Location/Qualifiers

FEATURES

source

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 male brains, age range 23-27 yo. Library is oligo-dt
 primed and directionally cloned (EcoRV site is destroyed
 upon cloning). Average insert size 1.5 kb, insert size
 range 1-3 kb. Library is normalized and enriched for
 full-length clones and was constructed by C. Gruber
 (Invitrogen). Research Genetics tracking code 019. Note:
 this is a NIH_MGC library."

ORIGIN

Alignment Scores:

Pred. No.: 1,72e-58 Length: 653
 Score: 602.00 Matches: 125
 Percent Similarity: 100.00% Conservative: 2
 Best Local Similarity: 98.43% Mismatches: 0
 Query Match: 98.69% Indels: 0
 DB: 4 Gaps: 0
 US-09-017-715a-2 (1-127) x B1757131 (1-653)

QY 1 MetAaPvAlPheLySgLyPheSerIleAlaLySgLyVAlVaIgIyAlaVaIgiu 20
 DB 57 ATGGATGCTTCAAGAGGCGCTTCTCCATCGCCAGAGGCGCTGCGGCGGAGAA 116
 QY 21 LysThrLySGInGlyValThrGluAlaIaGluLyThrLySGInGlyValMetTyVal 40
 DB 117 AAGACCAACAGGCGGTGACGAGACAGCTGAGAAACCAAGAGGCGGTCTATGTG 176
 QY 41 G1yAlaLyThrLySGInuSnVAlValInSerValThrSerValAlaGluLyThrLyS 60
 DB 177 GGAGCCAAAGACCAAGAGATGTTGTACAGAGCGTCACTGCGCCAGAAAGACCAAG 236
 QY 61 GluGlnAlaAsnAlaValSerLySAlaValSerValAsnThrValAlaThrLyS 80
 DB 237 GAGCAGGCGCAAGCGCTGAGCGGTGAGAGCGTCAACCTGTGGCCACCAAG 296
 QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValValArgLySGInuPleu 100
 DB 297 ACCGTGAGAGGCGGAGAAACATCGCGTCACTCCGGGCTGTGCGCAAGAGCACTTG 356
 QY 101 ArgProSerAlaProGInGInGluGlyGluAlaSerLySGInuGluGluValAlaGlu 120
 DB 357 AGGCCATCTGCCCCCAACAGAGGCGTGAAGCATCCAAAGAAAGAGAAAGTGCAGAG 416
 QY 121 GluAlaGlnSerGlyGlyasp 127
 DB 417 GAGGCCCAAGTGGGGAGAC 437
 RESULT 6
 B1836596 751 bp mRNA linear EST 04-OCT-2001
 LOCUS B1836596 603089575F1 NIH_MGC_120 Homo sapiens cDNA clone IMAGE:5228538 5',
 DEFINITION mRNA sequence.
 ACCESSION B1836596
 VERSION B1836596.1 GI:15948146

KEYWORDS

EST.
 Homo sapiens (human)

SOURCE

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 751)

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

NIH-MGC <http://mhc.nci.nih.gov/>.
 National Institutes of Health, Mammalian Gene Collection (MGC)
 Unpublished (1999)
 Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-remail.nih.gov
 Tissue Procurement: Life Technologies, Inc.
 cDNA Library Preparation: Life Technologies, Inc.
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
 Plate: L1M11575 row: e column: 19
 High quality sequence stop: 703.
 Location/Qualifiers

FEATURES

source

1..751
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 /clone_lib="NIH_MGC_120"
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 source anonymous pool of spleen and pancreas from 28 yo
 male. Library is oligo-dt primed and directionally cloned
 (EcoRV site is destroyed upon cloning). Average insert
 size 1.5 kb, insert size range 1-2.5 kb. Library is
 normalized and enriched for full-length clones and was
 constructed by C. Gruber (Invitrogen). Research Genetics
 tracking code 025. Note: this is a NIH_MGC library."

ORIGIN

Alignment Scores:

Pred. No.: 2,06e-58 Length: 751
 Score: 602.00 Matches: 125
 Percent Similarity: 100.00% Conservative: 2
 Best Local Similarity: 98.43% Mismatches: 0
 Query Match: 98.69% Indels: 0
 DB: 4 Gaps: 0
 US-09-017-715a-2 (1-127) x B1836596 (1-751)

QY 1 MetAaPvAlPheLySgLyPheSerIleAlaLySgLyVAlVaIgIyAlaVaIgiu 20
 DB 96 ATGGATGCTTCAAGAGGCGCTTCTCCATCGCCAGAGGCGCTGCGGCGGAGAA 155
 QY 21 LysThrLySGInGlyValThrGluAlaIaGluLyThrLySGInGlyValMetTyVal 40
 DB 156 AAGACCAACAGGCGGTGACGAGACAGCTGAGAAACCAAGAGGCGGTCTATGTG 215
 QY 41 G1yAlaLyThrLySGInuSnVAlValInSerValThrSerValAlaGluLyThrLyS 60
 DB 216 GGAGCCAAAGACCAAGAGATGTTGTACAGAGCGTCACTGCGCCAGAAAGACCAAG 275
 QY 61 GluGlnAlaAsnAlaValSerLySAlaValSerValAsnThrValAlaThrLyS 80
 DB 276 GAGCAGGCGCAAGCGCTGAGCGGTGAGAGCGTCAACCTGTGGCCACCAAG 335
 QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValValArgLySGInuPleu 100
 DB 336 ACCGTGAGAGGCGGAGAAACATCGCGTCACTCCGGGCTGTGCGCAAGAGCACTTG 395
 QY 101 ArgProSerAlaProGInGInGluGlyGluAlaSerLySGInuGluGluValAlaGlu 120
 DB 396 AGGCCATCTGCCCCCAACAGAGGCGTGAAGCATCCAAAGAAAGAGAAAGTGCAGAG 455

QY 121 Glu1a1aInserg1ygl1asp 127
 DB 456 GAGGCCAGAGTGGGGAGAGAC 476
 RESULT 7
 LOCUS B0893395
 DEFINITION AGENCOURT 8121065 Lupeki dorsal root ganglion Homo sapiens cDNA
 clone IMAGE:6178582 5', mRNA sequence.
 ACCESSION B0893395
 VERSION B0893395.1 GI:22285409
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 781)
 NIH-MGC http://mgi.nci.nih.gov/.
 AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-remail.nih.gov
 Tissue Procurement: Dr. James R. Lupski
 CDNA Library Preparation: Life Technologies, Inc.
 CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
 DNA Sequencing by: Agencourt Bioscience Corporation
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LNL at:
 http://image.llnl.gov
 Plate: LLM13558 row: F column: 23
 High quality sequence stop: 436.
 FEATURES
 source
 1. 781
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:6178582"
 /sex="male"
 /tissue_type="dorsal root ganglia"
 /dev_stage="adult, 36 yr"
 /lab_host="DH10B"
 /clone_lib="Lupeki dorsal root ganglion"
 /note="Vector: pCMV-SPORT6 (Life Technologies); Site 1:
 NotI; Site 2: SalI; cDNA made by oligo-dT priming.
 Directionally cloned using the following adaptors:
 5'-TCGACCCAGCGGTCG-3' and
 5'-GACTAGCTTCAGATCGGAGCGGCCGCTT(15)-3'. Size selected >
 1 kb for average insert length 1.7 kb. This is a primary
 library, non-amplified. Library constructed by Life
 Technologies and donated by J. Lupski, M.D./Ph.D. (Baylor
 College of Medicine) and is available through Life
 Technologies."
 ORIGIN
 Alignment Scores:
 Pred. No.: 2,16e-58 Length: 781
 Score: 602.00 Matches: 125
 Percent Similarity: 100.00% Conservative: 2
 Best Local Similarity: 98.43% Mismatches: 0
 Query Match: 98.69% Indels: 0
 Gaps: 0
 DB: 5
 US-09-017-715A-2 (1-127) x B0893395 (1-781)
 QY 1 MetAspValPheIyLysGlyPheSerIleAlaIyLysGlyValAlaIyAlaIyGlu 20
 DB 71 ATGATGCTCTTCAAGAGGCGCTTCTCCATCGCCAGAGAGGCGGCGGCGGTGAA 130
 QY 21 LysThrIyGlnGlyValIThrGluAlaIyGluIyLysGlyGluIyValMetIyVal 40
 DB 131 AAGACCAAGAGGCGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 190
 QY 41 G1yAlaIyThrIyGluAanValIyGlnSerValIyThrSerValIyAlaIyGluIyThrIy 60

DB 191 GAGGCCAGAGCCAGAGGAATGTTTTCACAGGCTGACTCAGTGGCCGAGAGACCAAG 250
 QY 61 Glu1a1aInserg1ygl1asp 127
 DB 251 GAGGCCAGAGCCAGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 310
 QY 81 ThrValGluIyGluAanValIyGlnSerValIyThrSerValIyAlaIyGluIyThrIy 100
 DB 311 ACCGTGAGAGGCGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 370
 QY 101 ArgProSerAlaProGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGln 120
 DB 371 AGGCGATCTGCCCCCAGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 430
 QY 121 Glu1a1aInserg1ygl1asp 127
 DB 431 GAGGCCAGAGTGGGGAGAGAC 451
 RESULT 8
 DB B0921124
 LOCUS B0921124
 DEFINITION AGENCOURT 6633376 NIH_MGC_115 Homo sapiens cDNA clone IMAGE:5752462
 5', mRNA sequence.
 ACCESSION B0921124
 VERSION B0921124.1 GI:19371503
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 809)
 NIH-MGC http://mgi.nci.nih.gov/.
 AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-remail.nih.gov
 Tissue Procurement: Life Technologies, Inc.
 CDNA Library Preparation: Life Technologies, Inc.
 CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
 DNA Sequencing by: Agencourt Bioscience Corporation
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LNL at:
 http://image.llnl.gov
 Plate: LLM12786 row: K column: 23
 High quality sequence stop: 738.
 FEATURES
 source
 1. 809
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:5752462"
 /lab_host="DH10B"
 /clone_lib="NIH_MGC_115"
 /note="Organ: pooled brain, lung, testis; Vector:
 pCMV-SPORT6; Site 1: NotI; Site 2: EcoRV (destroyed); RNA
 source anonymous pool of 6 male brains, age range 23-27; 1
 male lung, age 27; and 1 male testis, age 69. Library is
 oligo-dT primed and directionally cloned (EcoRV site is
 destroyed upon cloning). Average insert size 1.8 kb,
 insert size range 1-3 kb. Library is normalized and
 enriched for full-length clones and was constructed by C.
 Gruber (Invitrogen). Research Genetics tracking code
 021. Note: this is a NIH_MGC Library."
 ORIGIN
 Alignment Scores:
 Pred. No.: 2,26e-58 Length: 809
 Score: 602.00 Matches: 125
 Percent Similarity: 100.00% Conservative: 2
 Best Local Similarity: 98.43% Mismatches: 0
 Query Match: 98.69% Indels: 0
 DB: 5
 Gaps: 0

US-09-017-715A-2 (1-127) x BM921124 (1-809)

QY 1 MetaspvalPheLysGlyPheSerIleAlaLysGlyValGlyAlaValGlu 20
DB 114 ATGATGTCCTTCAAGAGGGCTTCTCCATCGCCCAAGAGGGCGTGTGGTGGTGGAA 173

QY 21 LysThrLysGlnGlyValThrGlnAlaGlnLysThrLysGlnGlyValMetYrVal 40
DB 174 AAGACCAAGAGGGGTGAGGAGAGCTTGAGAGACCAAGAGGGGTCTATGTAATG 233

QY 41 G1yAlaLysThrLysGlnLysValValGlnSerValThrSerValAlaGlnLysThrLys 60
DB 234 GGAGCCAAACCAAGAGAGATGTTGACAGAGGTGACCTCAGTGGCCGAGAAAGCAAG 293

QY 61 G1uGlnAlaAsnAlaValSerLysAlaValSerSerValAsnThrValAlaThrLys 80
DB 294 GAGCAGGCCCAAGCGCTGAGCGAGCTGTGTGAGCAGCGTCAACACTGTGGCCACCAAG 353

QY 81 ThrValGlnGlnAlaGlnLysValAlaValThrSerGlyValAlaArgLysGlnLysPleu 100
DB 354 ACCGTGAGAGAGCGAGAAACATCGCGTCACTCCGGGGTGTGTGCGCAAGAGGACTTG 413

QY 101 ArgProSerAlaProGlnGlnGlnGlnGlnLysGlnLysGlnLysGlnLysValAlaGln 120
DB 414 AAGCCATCTGCCCGCCCAACAGAGAGGTGAGGATCCAAAGAGAGAGAGAGTGGCAGAG 473

QY 121 G1uAlaGlnSerGlyGlyAsp 127
DB 474 GAGGCCCAAGTGTGGGAGAGAC 494

RESULT 9
BQ882072 855 bp mRNA linear EST 16-ANG-2002

LOCUS BQ882072
DEFINITION AGENCOURT 8586140 lupski_sympathetic_trunk Homo sapiens cDNA clone
IMAGE:6195522 5', mRNA sequence.
BQ882072
BO882072.1 GI:22274080

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
EST.
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 855)
NIH-MGC http://mgi.nci.nih.gov/
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgaabs-r@mail.nih.gov
Tissue Procurement: Dr. James R. Lupski
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLM13602 row: h column: 19
High quality sequence stop: 667.

FEATURES
source
Location/Qualifiers
1. 855
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:6195522"
/sex="male"
/tissue_type="sympathetic trunk"
/dev_stage="adult, 16 yr"
/lab_host="DH10B"
/clone_lib="lupski_sympathetic trunk"
/notes="Vector: pCMV-SPORT6 (Life Technologies); Site 1:
NotI; Site 2: SalI; cDNA made by oligo-dT priming.
Directionally cloned using the following adaptors:
5'-TCGACCCACGCGTCG-3' and

5'-GACTAGTCTTAGATCGAGCGGCCCT(15)-3'. Size selected >
1 kb for average insert length 1.9 kb. This is a primary
library non-amplified. Library constructed by Life
Technologies and donated by J. Lupski, M.D./Ph.D. (Baylor
College of Medicine); available through Life
Technologies."

ORIGIN

Alignment Scores:
Pred. No.: 2,436-58 Length: 855
Score: 602.00 Matches: 125
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 98.43% Mismatches: 0
Query Match: 98.69% Indels: 0
DB: Gaps: 0

US-09-017-715A-2 (1-127) x BQ882072 (1-855)

QY 1 MetaspvalPheLysGlyPheSerIleAlaLysGlyValGlyAlaValGlu 20
DB 94 ATGATGTCCTTCAAGAGGGCTTCTCCATCGCCCAAGAGGGGTGTGGTGGTGGAA 153

QY 21 LysThrLysGlnGlyValThrGlnAlaGlnLysThrLysGlnGlyValMetYrVal 40
DB 154 AAGACCAAGAGGGGTGAGGAGAGCTTGAGAGACCAAGAGGGGTCTATGTAATG 213

QY 41 G1yAlaLysThrLysGlnLysValValGlnSerValThrSerValAlaGlnLysThrLys 60
DB 214 GAGCCAAAGACCAAGAGAGATGTTGACAGAGGTGACCTCAGTGGCCGAGAAAGCAAG 273

QY 61 G1uGlnAlaAsnAlaValSerLysAlaValSerSerValAsnThrValAlaThrLys 80
DB 274 GAGCAGGCCCAAGCGCTGAGCGAGCTGTGTGAGCAGCGTCAACACTGTGGCCACCAAG 333

QY 81 ThrValGlnGlnAlaGlnLysValAlaValThrSerGlyValAlaArgLysGlnLysPleu 100
DB 334 ACCGTGAGAGAGCGAGAAACATCGCGTCACTCCGGGGTGTGTGCGCAAGAGAGACTTG 393

QY 101 ArgProSerAlaProGlnGlnGlnGlnGlnLysGlnLysGlnLysGlnLysValAlaGln 120
DB 394 AAGCCATCTGCCCGCCCAACAGAGAGGTGAGGATCCAAAGAGAGAGAGTGGCAGAG 453

QY 121 G1uAlaGlnSerGlyGlyAsp 127
DB 454 GAGGCCCAAGTGTGGGAGAGAC 474

RESULT 10
BQ439430 884 bp mRNA linear EST 24-MAY-2002

LOCUS BQ439430
DEFINITION AGENCOURT 7911914 NIH_MGC_68 Homo sapiens cDNA clone IMAGE:6010114
5', mRNA sequence.
BQ439430
BQ439430.1 GI:21178506

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
EST.
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 884)
NIH-MGC http://mgi.nci.nih.gov/
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgaabs-r@mail.nih.gov
Tissue Procurement: DCTD/DTF/Gazdar
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLM13197 row: k column: 11
High quality sequence stop: 559.

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

FEATURES

Source

```

Location/Qualifiers
1. .884
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:6010114"
/tissue_type="large cell carcinoma"
/lab_host="DH10B (phage-resistant)"
/clone_1fb="NH MCC_68"
/note="Organ: lung; Vector: pCMV-SPORT6; Site: 1: NotI; Site: 2: SalI; Cloned unidirectionally. Primer: oligo dT. Average insert size 1.8 kb. Library constructed by Life Technologies."

```

Alignment Scores:

Pred. No.:	2.58e+8	Length:	884
Score:	602.00	Matches:	125
Percent Similarity:	100.00%	Conservative:	2
Best Local Similarity:	98.43%	Mismatches:	0
Query Match:	98.69%	Indels:	0
DB:	5	Gaps:	0

AUTHORS NIH-MGC <http://mhc.nci.nih.gov/>.
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-r@mail.nih.gov
 Tissue Procurement: ATCC
 cDNA Library Preparation: Life Technologies, Inc.
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
 Plate: LHM10366 row: k column: 14
 High quality sequence stop: 681.

FEATURES
 source
 1..903
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:450613"
 /tissue_type="transitional cell papilloma, cell line"
 /lab_host="DH10B (phage-resistant)"
 /clone_lib="NIH_MGC_93"
 /note="Organ: bladder; Vector: pCMV-SPORT6; Site_1: NotI; Site_2: SalI; Cloned unidirectionally; oligo-dT primed. Average insert size 1.7 kb. Library enriched for full-length clones and constructed by Life Technologies. Note: this is a NIH_MGC Library."

ORIGIN

Alignment Scores:

Pred. No.:	2,61e-58	Length:	903
Score:	602.00	Matches:	125
Percent Similarity:	100.00%	Conservative:	2
Best Local Similarity:	98.43%	Mismatches:	0
Query Match:	98.69%	Indels:	0
DB:	4	Gaps:	0

US-09-017-715A-2 (1-127) x BG286466 (1-903)

```

QY      1 MetAspValPheIysLysGlyPheSerIleAlaLysGlyValValGlu 20
DB      87 ATGAGATGCTTCACAGAGGGCTTCATCGCCAGAGGGCGTGGTGGTGAA 146
QY      21 LysThrIysGlnGlyValThrGluAlaGluLysThrIysGlnGlyValMetTyrVal 40
DB      147 AAGACCAACAGAGGGGTGAACGAGACGCTGAGAACACAGAGGGGTGATGATG 206
QY      41 GAlAAlaLysThrIysGluLysValValGlnSerValThrSerValAlaGluLysThrIys 60
DB      207 GAGGCCAACACAGAGGATGTTGACAGAGGCTACCTGCGCCAGAACACAG 266
QY      61 GluGlnAlaAsnAlaValSerLysValValValSerSerValAsnThrValAlaThrLys 80
DB      267 GAGCAGGCCAACGCCCTGAGCGAGGCTGTGTGAGAGCGTCAACACTGTGGCCACCAAG 326
QY      81 ThrValGluGluAlaGluLysIleAlaValThrSerGlyValValAlaArgLysGluAspLeu 100
DB      327 ACCGTGAGAGAGCGGAGAACATCGCGGTCACTCGGGGTGTGTGCAAGAGGAGCTTG 386
QY      101 ArgProSerAlaProGlnGlnGlnGlyGluAlaSerLysGluLysGluGluValAlaGlu 120
DB      387 AGGCCATCTGCCCCCAACAGAGGGGTGAGGCTCCAAAGAGAAAGAGAGTGGCAGAG 446
QY      121 GluAlaGlnSerGlyGlyAsp 127
DB      447 GAGGCCACAGATGGGGAGAC 467

RESULT 13
LOCUS   B1548891          558 bp          mRNA          linear      EST 05-SEP-2001
DEFINITION 603189023F1 NIH_MGC_95 Homo sapiens cDNA clone IMAGE:5260589 5',
          mRNA sequence.

```

ACCESSION B1548891
 VERSION B1548891.1 GI:15436203
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens

REFERENCE
 AUTHORS Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 JOURNAL Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 558)
 NIH-MGC <http://mhc.nci.nih.gov/>.
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-r@mail.nih.gov
 Tissue Procurement: Miklos Palkevics, M.D., Ph.D.
 cDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki
 Toshiyuki and Piero Carninci (RIKEN)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
 Plate: LHM1656 row: m column: 06
 High quality sequence stop: 558.

FEATURES
 source
 1..558
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:5260589"
 /tissue_type="hippocampus"
 /lab_host="DH10B"
 /clone_lib="NIH_MGC_95"
 /note="Organ: brain; Vector: pBluescriptPR (modified pBluescript KS+); Site_1: BamHI; Site_2: SalI-XhoI (96cga); Oligo-dT primed using primer 5'-TTTTTCTTTTCTTTTCTTT-3', size-selected for average insert size 2.5 kb and normalized to ROP 5. This is a primary library enriched for full-length clones and constructed using the Cap-trapper method (Carninci, in preparation). Library constructed by M. Brownstein (NHGRI), National Institutes of Health. Note: this is a NIH_MGC Library."

ORIGIN

Alignment Scores:

Pred. No.:	8,82e-58	Length:	558
Score:	595.00	Matches:	124
Percent Similarity:	99.21%	Conservative:	2
Best Local Similarity:	97.64%	Mismatches:	1
Query Match:	97.54%	Indels:	0
DB:	4	Gaps:	0

US-09-017-715A-2 (1-127) x B1548891 (1-558)

```

QY      1 MetAspValPheIysLysGlyPheSerIleAlaLysGlyValValGlu 20
DB      130 ATGAGATGCTTCACAGAGGGCTTCATCGCCAGAGAGGGCGTGGTGCGGAGAA 189
QY      21 LysThrIysGlnGlyValThrGluAlaGluLysThrIysGlnGlyValMetTyrVal 40
DB      190 AAGACCAACAGAGGGGTGAACAGACGCTGAGAACACAGAGGGGTGATGATG 249
QY      41 GAlAAlaLysThrIysGluLysValValGlnSerValThrSerValAlaGluLysThrIys 60
DB      250 GAGGCCAACACAGAGGATGTTGACAGAGCGTCACTGCGCCAGAACACCAAG 309
QY      61 GluGlnAlaAsnAlaValSerLysValValValSerSerValAsnThrValAlaThrLys 80
DB      310 GAGCAGGCCAACGCCGTGAGCGAGGCTGTGTGAGAGCGTCAACACTGTGGCCACCAAG 369
QY      81 ThrValGluGluAlaGluLysIleAlaValThrSerGlyValValAlaArgLysGluAspLeu 100
DB      370 ACCGTGAGAGAGCGGAGAACATCGCGGTCACTCGGGGTGTGTGCAAGAGGAGCTTG 429

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Qy 101 ArgProSerAlaProGlnGlnGlyGluAlaSerIlyGlyGluGluValAlaGlu 120
 Db 430 AGGCATCTGCCCCCAAGAGAGGTGTGGCATCCAAAGAGAAAGAGAGTGGCAGAG 489
 Qy 121 GluAlaGlnSerGlyGlyAap 127
 Db 490 GAGGCCAGAGTGGGGAGAGC 510

RESULT 14
 BG708703 568 bp mRNA linear EST 07-MAY-2001
 LOCUS 602674249F1 NIH_MGC_96 Homo sapiens cDNA clone IMAGE:4796820 5',
 DEFINITION mRNA sequence.
 BG708703
 VERSION BG708703.1 GI:13986308
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 568)
 AUTHORS NIH-MGC http://mgs.nci.nih.gov/
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgaabs-remail.nih.gov
 Tissue Procurement: Miklos Palokovits, M.D., Ph.D.
 cDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki
 Toshiyuki and Piero Carninci (RIKEN)
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LNL at:
 http://image.llnl.gov
 Plate: LLM10682 row: a column: 13
 High quality sequence stop: 566.
 Location/Qualifiers
 1. 568
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:4796820"
 /tissue_type="hypothalamus"
 /lab_host="DH10B"
 /clone_lib="NIH_MGC_96"
 /note="Organ: brain; Vector: pBluescriptR (modified
 pBluescript KS+); Site: 1: BamHI; Site 2: SalI-XhoI
 (gfcgag); Oligo-dT primed using primer
 5'-TTTTTTTTTTTTTN-3', size-selected for average
 insert size 2.3 kb and normalized to ROP 5. This is a
 primary library enriched for full-length clones and
 constructed using the Cap-trapper method (Carninci, in
 preparation). Library constructed by M. Brownstein
 (NIH/NHGRI, National Institutes of Health). Note: this is
 a NIH_MGC Library."

ORIGIN
 Alignment Scores:
 Pred. No.: 9.02e-58 Length: 568
 Score: 595.00 Matches: 124
 Percent Similarity: 99.21% Conservative: 2
 Best Local Similarity: 97.64% Mismatches: 1
 Query Match: 97.54% Indels: 0
 DB: 4 Gaps: 0

US-09-017-715A-2 (1-127) x BG708703 (1-568)

Qy 1 MetAspAlaPheIlyGlyGlyPheSerIAlaIlyGlyGlyValAlaGlyAlaValGlu 20
 Db 108 ATGAGATGCTTCAAGAGGCTTCTCCATGCGCAAGAGGCGGTGGGTGGCGTGGAA 167

Qy 21 LyeThrIlyGlnGlnGlyValThrGluAlaAlaGluIlyThrIlyGlnGluGlyValMetCTYVal 40

Db 168 AAGACCAAGAGGGGGGTGACGGAAGCCTGAGAGGCCAAGAGGGGCTCATGTATGTG 227
 Qy 41 GlYAlaIlyThrIlyGlnGlnValAlaGlnSerValThrSerValAlaGluIlyThrIly 60
 Db 228 GAGGCCAAGACCAAGAGGAATGTTGTACAGAGCTGACTCAGTGGCGGAGAACCAAG 287
 Qy 61 GluGlnAlaSerAlaValSerIlyAlaValIlySerSerValIleThrValAlaThrIly 80
 Db 288 GAGAGGCCAAGCGGTGAGCGGTGTGTAGCAGCGCTCAACACTGTGGCCACCAAG 347
 Qy 81 ThrValGlnGluAlaGlnIleAlaValThrSerGlyValAlaGlyGlyGluAspLeu 100
 Db 348 ACCGTGAGAGAGCGGAGAACATCGCGTCACTCCGGGTGTGGCCAGAGACTTG 407
 Qy 101 ArgProSerAlaProGlnGlnGlyGluAlaSerIlyGlyGluGluValAlaGlu 120
 Db 408 AGGCATCTGCCCCCAAGAGAGGTGTGGCATCCAAAGAGAAAGAGAGTGGCAGAG 467
 Qy 121 GluAlaGlnSerGlyGlyAap 127
 Db 468 GAGGCCAGAGTGGGGAGAGC 488

RESULT 15
 BG707764 617 bp mRNA linear EST 07-MAY-2001
 LOCUS 602671103F1 NIH_MGC_96 Homo sapiens cDNA clone IMAGE:4793833 5',
 DEFINITION mRNA sequence.
 BG707764
 VERSION BG707764.1 GI:13984439
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 617)
 AUTHORS NIH-MGC http://mgs.nci.nih.gov/
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgaabs-remail.nih.gov
 Tissue Procurement: Miklos Palokovits, M.D., Ph.D.
 cDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki
 Toshiyuki and Piero Carninci (RIKEN)
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LNL at:
 http://image.llnl.gov
 Plate: LLM10674 row: e column: 02
 High quality sequence stop: 617.
 Location/Qualifiers
 1. 617
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:4793833"
 /tissue_type="hypothalamus"
 /lab_host="DH10B"
 /clone_lib="NIH_MGC_96"
 /note="Organ: brain; Vector: pBluescriptR (modified
 pBluescript KS+); Site: 1: BamHI; Site 2: SalI-XhoI
 (gfcgag); Oligo-dT primed using primer
 5'-TTTTTTTTTTTTTN-3', size-selected for average
 insert size 2.3 kb and normalized to ROP 5. This is a
 primary library enriched for full-length clones and
 constructed using the Cap-trapper method (Carninci, in
 preparation). Library constructed by M. Brownstein
 (NIH/NHGRI, National Institutes of Health). Note: this is
 a NIH_MGC Library."

ORIGIN
 Alignment Scores:

Pred. No.: 1e-57 Length: 617
Score: 595.00 Matches: 124
Percent Similarity: 99.21% Conservative: 2
Best Local Similarity: 97.64% Mismatches: 1
Query Match: 97.54% Indels: 0
DB: 4 Gaps: 0

US-09-017-715a-2 (1-127) x BG707764 (1-617)

```
QY      1 MetAEPValPheLyLeLySerIleAlaLyLeLyGlyValValGlyAlaValGlu 20
DB      58 ATGGATGCTTCAGAGAGGCTTCTCCATCGCCAGAGGGCGGTGGTGGCGGAGAA 117
QY      21 LysThrLySGInGlyValThrGluAlaGluLyThrLySGInGlyValMetTyrVal 40
DB      118 AAGACCAAGCAGGGGGGTGCGAGAGCACTGAGAGAACCAAGAGGGGGTCAATGTA 177
QY      41 GlyAlaLySThrLySGInuAsnValGlnSerValThrSerValAlaGluLySThrLy 60
DB      178 GGAGCCAGAGCAAGAGAAATGTTGTACAGAGCGGTGACCTCACTGCGCAGAAAGACCAAG 237
QY      61 GluGlnAlaAsnAlaValSerLySAlaValSerSerValAsnThrValAlaThrLy 80
DB      238 GAGCAGGCGCAAGCGGTGAGCGAGCTGTGTGAGCAGCGTCAACACTGTGGCCACCAAG 297
QY      81 ThrValGluGluValaGluAsnIleAlaValThrSerGlyValValArgLySGInuAspLeu 100
DB      298 ACCGTGGAGAGGCGGAGAAACATCGCGGTCACTCCGGGGTGTGGCAAGAGAGACTTG 357
QY      101 ArgProSerAlaProGInGInGInGlyGluAlaSerLySGInuLySGInuValAlaGlu 120
DB      358 AGGCCATGTGCCCGCCCAAGAGAGGGGTGTGCATCCAAAGAGAAAGAGAGAGTGGCAGAG 417
QY      121 GluAlaGlnSerGlyGly*Sp 127
DB      418 GAGGCCAGAGTGGGGGAGAC 438
```

Search completed: May 4, 2005, 13:46:32
Job time : 5788.07 secs

GenCore version 5.1.6
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OM protein - nucleic search, using frame_plus_p2n model

Run on: May 4, 2005, 09:07:52 ; Search time 58.4698 Seconds

(without alignments)

809.955 Million cell updates/sec

Title: US-09-017-715A-2_COPY_120_127

Perfect score: 41

1 EBAQSGSD 8

Scoring table:

BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Command line parameters:

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-O=/cgn2.1/USPTO.spool.h/US09017175/runat.04052005.100743.25600/app.query.fasta.1.661
-DB=N_Geneseq -OFMT=fastcap -SUFFIX=ring -MINMATCH=0.1 -LOOFC=0 -LOOPEXT=0
-UNITS=bits -START=1 -END=1 -MATRIX=blomum62 -TRANS=human40.cdi -LIST=45
-DOCALLIGN=200 -THR_SCORE=pct -THR_MAX=100 -THR_MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTFMT=pct -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=200000000
-USBR=US09017175 @CGN 1.1 703 @runat.04052005.100743.25600 -NCPD=6 -ICPU=3
-NO_MMAP -LARGEQUERRY -NEG_SCORES=0 -WAIT -DSPELOCK=100 -LONGLOG
-DEV TIMEOUT=120 -MAIN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database : N_Geneseq_16Dec04:*

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Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result	Query	Match	Length	DB	ID	Description
No.	Score					
1	41	100.0	478	3	AAF21785	Aaf21785 Human bre
2	41	100.0	479	9	ACH15493	Ach15493 Human adu
3	41	100.0	488	12	ADM66887	Adm66887 Human hom
4	41	100.0	550	2	AAV42669	Aav42669 Human bre
5	41	100.0	550	3	AAA39470	Aaa39470 Human HBG

6	41	100.0	550	6	ABL63343	ABL63343 Breast ca
7	41	100.0	550	6	ABV73813	Abv73813 Human gam
8	41	100.0	550	6	ABV73915	Abv73915 Human gam
9	41	100.0	550	10	AAD63568	Aad63568 Human amy
10	41	100.0	550	10	ADG47636	Adg47636 Human amy
11	41	100.0	720	2	AAx29997	Aax29997 Human per
12	41	100.0	720	6	ABs76519	AbS76519 cDNA enco
13	41	100.0	720	10	ADs43864	AdS43864 Human SNC
14	41	100.0	720	12	ADs43432	AdS43432 Human SNC
15	41	100.0	796	3	AAI93778	Aai93778 Human pol
16	41	100.0	796	3	AAI93784	Aai93784 Human pol
17	41	100.0	990	13	ADR98806	AdR98806 Lung spec
18	38	92.7	1125	13	ADs57612	AdS57612 Bacterial
19	38	92.7	110000	11	ADM27081_11	Continuation (12 o
20	37	90.2	1275	4	ABL04107	ABL04107 Drosophi1
21	37	90.2	3275	4	ABL04106	ABL04106 Drosophi1
22	36	87.8	2116	10	ADB62945	AdB62945 Human CDN
23	36	87.8	3790	10	ADC39175	AdC39175 Novel hum
24	36	87.8	4062	13	ADQ89903	AdQ89903 Antagonis
25	36	87.8	4949	12	ADQ85477	AdQ85477 Human tum
26	36	87.8	4985	2	AAK00463	Aax00463 Human typ
27	36	87.8	4985	12	ADP21328	AdP21328 Gene ADcy
28	36	87.8	5054	10	ADF74204	AdF74204 Human nov
29	36	87.8	5236	10	ADC30279	AdC30279 Human nov
30	36	87.8	5372	12	ADQ25367	AdQ25367 Human sof
31	36	87.8	349980	6	ABO81847	AbO81847 Bifidobac
32	35	85.4	458	4	AAK76569	Aak76569 Human imm
33	35	85.4	458	4	AAK76570	Aak76570 Human imm
34	35	85.4	518	13	ACN48722	Acn48722 Cotton pr
35	35	85.4	547	12	ACH73352	Ach73352 Human gen
36	35	85.4	677	2	AAK04876	Aax04876 Human gam
37	35	85.4	918	13	ADT46476	AdT46476 Bacterial
38	35	85.4	1273	11	ACN91507	Acn91507 Breast ca
39	35	85.4	2143	10	ADC07759	AdC07759 Rice DNA
40	35	85.4	2154	10	ADC08252	AdC08252 Rice DNA
41	35	85.4	2251	10	AD160540	Ad160540 Secreved
42	35	85.4	3081	4	ABL27575	AB127575 Drosophi1
43	35	85.4	4259	10	ADs58369	AdS58369 Toxicity
44	35	85.4	4259	10	ADs52941	AdS52941 Primary r
45	35	85.4	4292	10	ADE71246	Ade71246 Novel hum

ALIGNMENTS

RESULT 1	AAF21785	standard; DNA; 478 BP.
ID	AAF21785	
XX	AAF21785;	
AC	27-MAR-2001	(first entry)
DT		
XX		
DE		Human breast and ovarian cancer associated antigen gene SEQ ID 172.
XX		
KW		Human; breast cancer; ovarian cancer; cytostatic; immunosuppressive;
KW		neurotrophic; neuroprotective; antiviral; antiallergic; hepatotropic;
KW		antidiabetic; antiinflammatory; antitumor; vulnerrary; anticonvulsant;
KW		antibacterial; antifungal; antiparasitic; cardiant; immune disorder;
KW		Addison's disease; allergy; autoimmune haemolytic anaemia;
KW		autoimmune thyroiditis; diabetes mellitus; Crohn's disease;
KW		multiple sclerosis; rheumatoid arthritis; ulcerative colitis;
KW		cardiovascular disorder; wound healing; neurological disease; ds.
XX		
OS		Homo sapiens.
PN		WO20005173-A1.
XX		
PD		21-SEP-2000.
XX		
PF		08-MAR-2000; 2000WO-US005881.
XX		
PR		12-MAR-1999; 99US-0124270P.
XX		

PA (HUMA-) HUMAN GENOME SCI INC.
XX Rosen CA, Ruben SM;
PI
XX MPI: 2000-611515/58.
XX P-PSDB; AAB58862.
DR
XX New human breast and ovarian cancer associated gene sequences and the
PT polypeptides encoded by these genes, useful in the prevention, treatment
PT and diagnosis of cancer, immune disorders, cardiovascular disorders and
PT neurological diseases.
PS Claim 1; Page 609; 1299pp; English.
XX
XX Sequences AAF21614 - AAF22031 represent DNA sequences encoding human
CC proteins AAB58711 - AAB59128. The DNA and protein sequences are
CC associated with breast and ovarian cancer. Included in the invention are
CC sequences AAF22032 - AAF22040 and AAB59129 which are used in the
CC isolation and characterization of the DNA and protein sequences of the
CC invention. The breast and ovarian cancer associated DNA, protein, agonist
CC or antagonist sequences exhibit cytostatic; immunosuppressive; neurotropic;
CC neuroprotective; antiviral; antiallergic; hepatotropic; antidiabetic;
CC antiinflammatory; antitumor; vulnery; anticonvulsant; antibacterial;
CC antifungal; antiparasitic and cardiac activity. The polynucleotide and
CC protein sequences are used in the diagnosis of cancer, particularly
CC breast and ovarian cancer. The nucleic acid sequences, proteins, agonists
CC and antagonists may also be used in the diagnosis, prevention and treatment
CC of immune disorders e.g. Addison's disease, allergies, autoimmune
CC haemolytic anemia, autoimmune thyroiditis, diabetes mellitus, Crohn's
CC disease, multiple sclerosis, rheumatoid arthritis and ulcerative colitis;
CC cardiovascular disorders such as myocardial ischaemia; wound healing;
CC neurological diseases such as cerebral anoxia and epilepsy; and
CC infectious diseases
SQ Sequence 478 BP; 118 A; 150 C; 113 G; 97 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 47.8 Length: 478
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 3 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x AAF21785 (1-478)

QY 1 GIUGUAlaGlnSerGlyGlyasp 8
DB 149 GAGGAGGCCCAAGATGTGGGAGAC 172

RESULT 2
ACH15493
ID ACH15493 standard; cDNA; 479 BP.
XX
XX ACH15493;
AC
XX
XX 13-OCT-2003 (first entry)
DT
XX
XX Human adult brain cDNA #2705.
DE
XX
XX Human; ss; sequencing by hybridisation; SBH; expressed sequence tag; EST;
KW genome mapping; biodiversity; genetic disorder.
XX
XX Homo sapiens.
OS
XX
XX US2003073623-A1.
PN
XX
XX 17-APR-2003.
PD
XX
XX 30-JUL-2001; 2001US-00918995.
PF
XX
XX 30-JUL-2001; 2001US-00918995.
PR
XX
XX

PA (DRMA/) DRMANAC R T.
PA (LABA/) LABAT I.
PA (STAC/) STACHE-CRAIN B.
PA (DICK/) DICKSON M C.
PA (JONE/) JONES L W.
XX
XX Drmanac RT, Labat I, Stache-Crain B, Dickson MC, Jones LW,
PI MPI: 2003-615964/58.
DR
XX
XX
PT New polynucleotide sequences obtained from various cDNA libraries, useful
PT as hybridization probes, as oligomers for PCR, for chromosome and gene
PT mapping, in the recombinant production of protein, or in generating
PT antisense DNA or RNA.
PS Claim 1; SEQ ID NO 2705; 44pp; English.
XX
XX The invention relates to an isolated polynucleotide comprising any one of
CC 38043 cDNA sequences, appearing as ACH12789-ACH50831, whose sequence was
CC determined by the technique of SBH (sequencing by hybridisation). Also
CC included is a purified polypeptide comprising a sequence corresponding to
CC a reading frame of the novel polynucleotide. The nucleic acid sequences
CC are useful in diagnostics as expressed sequence tags (EST) for
CC identifying expressed genes or for physical mapping of the human genome,
CC in forensics, in assessing biodiversity, or in identifying mutations
CC responsible for genetic disorders and other traits. The nucleotide
CC sequences are also useful as hybridisation probes, as oligomers for PCR,
CC for chromosome and gene mapping, in the recombinant production of
CC protein, or in generating antisense DNA or RNA. The purified polypeptide
CC is useful for generating antibodies specific for it. The present sequence
CC is one of the 38043 isolated cDNA/EST sequences. Note: The sequence data
CC for this patent did not form part of the printed specification, but was
CC obtained in electronic format directly from USPTO at
CC segdata.uspto.gov/sequence.html?docID=20030073623
SQ Sequence 479 BP; 120 A; 118 C; 178 G; 56 T; 0 U; 7 Other;

Alignment Scores:
Pred. No.: 47.9 Length: 479
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x ACH15493 (1-479)

QY 1 GIUGUAlaGlnSerGlyGlyasp 8
DB 371 GAGGAGGCCCAAGATGTGGGAGAC 394

RESULT 3
ADM66887
ID ADM66887 standard; DNA; 488 BP.
XX
XX ADM66887;
AC
XX
XX 03-JUN-2004 (first entry)
DT
XX
XX Human homologue of murine adipocyte specific gamma synuclein DNA Seq 20.
DE
XX
XX human; adipocyte specific; gene; ds; adipose tissue; anti-obesity;
KW high mobility group I-C protein; HMGI-C; obesity; lepin; ob; diabetes;
KW adipogenesis; hypertension; cardiovascular disease; anorectic;
KW antidiabetic; hypotensive; gamma synuclein.
XX
XX Homo sapiens.
OS
XX
XX WO2004011618-A2.
PN
XX
XX 05-FEB-2004.
PD
XX
XX 29-JUL-2003; 2003WO-US023684.
PF

XX 29-JUL-2002; 2002US-0398785P.
PR 12-JUN-2003; 2003US-0478206P.
XX (HMG-1) HMG-1 INC.
XX Chada K, Chouinard R, Aehar H, Sayed AMD;
PI WPI; 2004-143846/14.
XX P-PSDB; ADM67167.
XX
XX Identifying adipocyte specific genes, useful for treating obesity or
PT diabetes, and for identifying drug targets, by differential gene
PT expression analysis between adipose tissue or stromal vascular tissue of
PT mice of different genotypes.
PS Claim 11; SEQ ID NO 20; 91pp; English.
XX
XX This invention relates to a novel method for identifying genes that are
CC over-expressed in adipose tissue and as such it provides targets for anti-
CC obesity pharmaceutical compositions. Specifically, it refers to a high
CC mobility group I-C protein (HMG1-C) that is associated with obesity and
CC is epistatic to leptin, furthermore, it refers to the ob gene where an
CC autosomal recessive trait is linked to obesity and diabetes. The present
CC invention describes performing differential gene expression analysis
CC between the white adipose tissue (WAT) or stromal vascular tissue (SVT)
CC of any two different mice selected from a group consisting of wild-type,
CC HMG1-C^{-/-}, ob/ob, or HMG1-C^{-/-} ob/ob genotype mice. Accordingly, using
CC this method novel nucleotides and the encoded proteins thereof were
CC identified that are adipocyte specific, and as such can be used for
CC preventing adipogenesis, diagnosing and treating diabetes, obesity,
CC hypertension and cardiovascular disease, as well as screening for
CC compounds that can modulate or prevent adipogenesis and treat diabetes or
CC obesity. These compositions exhibit anorectic, antidiabetic and
CC hypopneumatic activities. This polynucleotide sequence is a human homologue
XX of a murine adipocyte specific DNA sequence of the invention.
SQ Sequence 488 BP; 127 A; 119 C; 176 G; 66 T; 0 U; 0 Other;
XX
XX Alignment Scores:
Pred. No.: 48.9 Length: 488
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 12 Gaps: 0
XX
US-09-017-715A-2_COPY_120_127 (1-8) x ADM66887 (1-488)
QY 1 GluGluAlaGlnSerGlyGlyAap 8
DB 369 GAGGAGGCCCGAGTGGGGGAGAC 392
XX
RESULT 4
AAV42669
ID AAV42669 standard; cDNA; 550 BP.
XX
AC AAV42669;
XX
DT 09-NOV-1998 (first entry)
XX
XX Human breast cancer specific gene 1 (BCSG1) cDNA.
DE
XX
XX Breast cancer specific gene 1; BCSG1; human; metastasis; diagnosis;
KW therapy; genetic marker; ds.
XX
XX Homo sapiens.
OS
XX
XX Key Location/Qualifiers
FH 12..395 /*tag= a
FT CDS
FT
FT
PN W09833915-A1.

XX 06-AUG-1998.
PD
XX 03-FEB-1998; 98MO-US001804.
PF
XX 03-FEB-1997; 97US-0037080P.
PR
XX (HUMA-) HUMAN GENOME SCI INC.
PA
XX J1 H, Rosen CA;
PI
XX WPI; 1998-446811/38.
DR
XX P-PSDB; AAW63123.
XX
XX New isolated human breast cancer specific gene - used to develop products
PT for the diagnosis, clinical management and treatment of breast cancer and
PT metastases.
PS Claim 4; Fig 1; 73pp; English.
XX
XX This cDNA clone corresponds to the transcript of the newly identified
CC human breast cancer specific gene 1 (BCSG1), and includes an open reading
CC frame for a 14.2 kDa protein (see AAW63123). It was isolated from a
CC breast cancer cDNA library following an EST search for novel genes
CC differentially expressed in breast cancer versus healthy breast tissue.
CC The clone is deposited at ATCC 97175 and ATCC 97856. A gradient and stage
CC -specific BCSG1 expression has been demonstrated from virtually no
CC detectable expression in normal or benign breast to low level and partial
CC expression in low grade in situ breast carcinoma and high expression in
CC infiltrating malignant breast carcinomas. BCSG1 is useful as a breast
CC cancer progression marker. Recombinant vectors and host cells useful for
CC recombinant production of BCSG1 polypeptides (including epitope-bearing
CC polypeptides) are provided. BCSG1 polynucleotides, polypeptides and
CC antibodies can be used for the detection of breast cancer cells or breast
CC cancer metastasis, and to develop methods for the clinical management and
XX treatment of breast cancer
SQ Sequence 550 BP; 132 A; 145 C; 192 G; 81 T; 0 U; 0 Other;
XX
XX Alignment Scores:
Pred. No.: 55.4 Length: 550
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 2 Gaps: 0
XX
US-09-017-715A-2_COPY_120_127 (1-8) x AAV42669 (1-550)
QY 1 GluGluAlaGlnSerGlyGlyAap 8
DB 369 GAGGAGGCCCGAGTGGGGGAGAC 392
XX
RESULT 5
AAA39470
ID AAA39470 standard; DNA; 550 BP.
XX
AC AAA39470;
XX
DT 24-AUG-2000 (first entry)
XX
XX Human HBGBA67A DNA.
DE
XX
XX Human; ADA2; cytosolic; gene therapy; treatment; cancer;
KW amyloid-like protein; ss.
XX
XX Homo sapiens.
OS
XX
XX Key Location/Qualifiers
FH 12..395 /*tag= a
FT CDS /*product= "HBGBA67"
FT
FT
FT
PN

PN US6054289-A.
XX
XX 25-APR-2000.
XX
XX 30-AUG-1996; 96US-00705771.
XX
XX 30-AUG-1995; 95US-0002993P.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Moore PA;
PI WPI; 2000-338491/29.
DR P-PSDB; AAY87779.
XX
XX New polynucleotide encoding human AD2 is useful for treating cancer and
PT for isolating cDNAs and genes having similar biological activity.
XX
XX Disclosure; Col 27-28; 54pp; English.
XX
XX This invention describes a novel polynucleotide (I) encoding human ADA2.
CC The products of the invention have cytoskeletal activity and can be used
CC for gene therapy. (I) is useful for treating cancer; as primers and
CC probes for isolating full length cDNA and genes having similar biological
CC activity. This sequence encodes a polypeptide derived from the human
CC HBGA67X clone which is an amyloid-like protein found in breast tissue
XX
SQ Sequence 550 BP; 132 A; 145 C; 192 G; 81 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 55.4 Length: 550
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservatve: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x AAA39470 (1-550)
QY 1 GIUGUAAGlnSerClyGlyAap 8
DB 369 GAGGAGGCCAGAGTGGGAGAC 392

RESULT 6
ABL63343
ID ABL63343 standard; DNA; 550 BP.
XX
XX ABL63343;
AC
XX
XX 15-MAY-2002 (first entry)
DT
XX
XX Breast cancer related gene sequence SEQ ID NO:1680.
DE
XX Human; cancer; colon; breast; ovary; oesophagus; kidney; thyroid;
KW stomach; lung; prostate; pancreas; carcinoma; antitumour; cancerous;
KM cytoskeletal; gene therapy; anti-neoplastic; Wilm's tumour; adenocarcinoma;
XX gene; ds.
XX
XX Homo sapiens.
OS
XX
XX WO200194629-A2.
PN
XX
XX 13-DEC-2001.
PD
XX
XX 30-MAY-2001; 2001WO-US010838.
PF
XX
XX 05-JUN-2000; 2000US-0209473P.
PR 05-JUN-2000; 2000US-0209531P.
PR 18-SEP-2000; 2000US-0233133P.
PR 18-SEP-2000; 2000US-023317P.
PR 20-SEP-2000; 2000US-0234009P.
PR 20-SEP-2000; 2000US-0234034P.
PR 20-SEP-2000; 2000US-0234052P.

PR 22-SEP-2000; 2000US-0234509P.
PR 22-SEP-2000; 2000US-0234567P.
PR 25-SEP-2000; 2000US-0234923P.
PR 25-SEP-2000; 2000US-0234924P.
PR 25-SEP-2000; 2000US-0235077P.
PR 25-SEP-2000; 2000US-0235082P.
PR 25-SEP-2000; 2000US-0235134P.
PR 25-SEP-2000; 2000US-0235280P.
PR 26-SEP-2000; 2000US-0235637P.
PR 26-SEP-2000; 2000US-0235638P.
PR 27-SEP-2000; 2000US-0235711P.
PR 27-SEP-2000; 2000US-0235720P.
PR 27-SEP-2000; 2000US-0235840P.
PR 27-SEP-2000; 2000US-0235863P.
PR 28-SEP-2000; 2000US-0236028P.
PR 28-SEP-2000; 2000US-0236033P.
PR 28-SEP-2000; 2000US-0236033P.
PR 28-SEP-2000; 2000US-0236033P.
PR 28-SEP-2000; 2000US-0236033P.
PR 28-SEP-2000; 2000US-0236109P.
PR 28-SEP-2000; 2000US-0236111P.
PR 28-SEP-2000; 2000US-0236111P.
PR 29-SEP-2000; 2000US-0236842P.
PR 29-SEP-2000; 2000US-0236891P.
PR 02-OCT-2000; 2000US-0237172P.
PR 02-OCT-2000; 2000US-0237173P.
PR 02-OCT-2000; 2000US-0237278P.
PR 02-OCT-2000; 2000US-0237294P.
PR 02-OCT-2000; 2000US-0237295P.
PR 02-OCT-2000; 2000US-0237315P.
PR 03-OCT-2000; 2000US-0237425P.
PR 03-OCT-2000; 2000US-0237598P.
PR 03-OCT-2000; 2000US-0237604P.
PR 03-OCT-2000; 2000US-0237606P.
PR 03-OCT-2000; 2000US-0237608P.
PR 01-NOV-2000; 2000US-0244867P.
PR 01-NOV-2000; 2000US-0245084P.
XX
XX (AVAL-) AVALON PHARM.
PA
XX Young PE, Augustus M, Carter KC, Ebner R, Endress G, Horrigan S;
PI Soppet DR, Weaver Z;
XX
XX WPI; 2002-188264/24.
DR
XX
XX Screening for anti-neoplastic agent involves exposing cells to a chemical
PT agent to be tested for anti-neoplastic activity, and determining a change
PT in expression of a gene of a signature gene set.
XX
XX Claim 1; SEQ ID NO 1680; 44pp; English.
XX
XX The present invention describes a method (M1) for screening for an anti-
CC neoplastic agent. The method involves exposing cells to a chemical agent
CC to be tested for anti-neoplastic activity, determining a change in
CC expression of at least one gene (I) of a signature gene set, where (I)
CC comprises a sequence (S) selected from 8447 sequences (given in ABL61664
CC to ABL70110), or is at least 95% identical to (S), where a change in
CC expression is indicative of anti-neoplastic activity. (I) has cytoskeletal
CC activity and can be used in gene therapy. M1 can be used for screening an
CC anti-neoplastic agent, and can be used for producing a product which is
CC the data collected with respect to the anti-neoplastic agent as a result
CC of M1, and the data is sufficient to convey the chemical structure and/or
CC properties of the agent. M1 can be used in the treatment of cancer such
CC as colon, breast, stomach, lung, thyroid, oesophageal, ovarian, kidney,
CC prostate or pancreatic cancer, adenocarcinoma, carcinoma, clear cell
CC cancer, infiltrating ductal cancer, infiltrating lobular cancer, squamous
CC cell carcinoma, neuroendocrine carcinoma, papillary carcinoma and Wilm's
CC tumour
XX
SQ Sequence 550 BP; 132 A; 145 C; 192 G; 81 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 55.4 Length: 550
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservatve: 0

Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-09-017-715a-2_COPY_120_127 (1-8) x ABV63343 (1-550)

Qy 1 GUGUUAAGInserGlyGlyAap 8
Db 369 GAGGAGCCCGAGAGTGGGGAGAC 392

RESULT 7
ABV73813
ID ABV73813 standard; cDNA; 550 BP.

XX ABV73813;

XX 08-JAN-2003 (first entry)

XX Human gamma-synuclein Glu110 variant gene.

XX Gamma-synuclein; human; single nucleotide polymorphism; SNP;

XX schizophrentia; neuroleptic; gene; ss.

XX Homo sapiens.

XX Key Location/Qualifiers
CDS 12..395

FT /*tag= a
FT /product= "Gamma-synuclein"
FT /transl_except= (pos:213..215,aa:Glu)
FT replace(340,T)
FT /*tag= b
FT /standard_name= "Single nucleotide polymorphism"

XX WO200275317-A2.

XX 26-SEP-2002.

XX 14-MAR-2002; 2002WO-EP002872.

XX 15-MAR-2001; 2001US-0276306P.

XX (NOVS) NOVARTIS AG.

XX (NOVS) NOVARTIS-ERFINDUNGEN VERW GES MBH.

XX (UYMA-) UNIV MARYLAND BALTIMORE.

XX Roberts RC, Van Oostrum J, Voshol J, Tamminga CA;

XX WPI; 2002-750574/81.

XX P-PsDB; ABP54932.

XX Screening for compounds for treating or interfering with the onset of
XX Schizophrenia Spectrum Disorders, by detecting interactions of candidate
XX compounds with the gamma-synuclein polypeptide.

XX Disclosure; Fig 1; 32pp; English.

XX The present sequence is that of cDNA encoding the Glu-110 isoform of
XX human gamma-synuclein. The invention relates to an isoform of gamma-
XX synuclein that is caused by an A/T single nucleotide polymorphism (SNP)
XX at position 329 of the gamma-synuclein coding sequence. This SNP causes a
XX glutamic acid to valine change at amino acid position 110 of gamma-
XX synuclein, and is associated with an increased susceptibility of
XX individuals to schizophrentia spectrum disorders (SSDs). This is the first
XX time that a genetic component of SSDs has been identified, and provides a
XX potential target for diagnosis and treatment of schizophrentia. Gamma-
XX synuclein polypeptides, especially those containing the E110V mutation,
XX are used in a claimed method of screening for compounds useful for the
XX treatment of SSDs, and gamma-synuclein expressing cells are used in a
XX claimed method of screening for agonist or antagonist compounds. An
XX oligonucleotide complementary to part of the gamma-synuclein coding
XX sequence is used for the discrimination of an SNP at position 329 of the
XX coding sequence. Gamma-synuclein polypeptides or polymucleotides are also

CC useful for the diagnosis of SSDs, or susceptibility to SSDs, e.g. by PCR
CC amplification of a polymucleotide encoding gamma-synuclein and analysis
CC of the occurrence of the SNP at position 329. A transgenic animal useful
CC for the study of SSDs is also claimed

XX Sequence 550 BP; 132 A; 145 C; 192 G; 81 T; 0 U; 0 Other;

XX Alignment Scores:

Pred. No.:	55.4	Length:	550
Score:	41.00	Matches:	8
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	6	Gaps:	0

US-09-017-715a-2_COPY_120_127 (1-8) x ABV73813 (1-550)

Qy 1 GUGUUAAGInserGlyGlyAap 8

Db 369 GAGGAGCCCGAGAGTGGGGAGAC 392

RESULT 8

ID ABV73915 standard; cDNA; 550 BP.

XX ABV73915;

XX 08-JAN-2003 (first entry)

XX Human gamma-synuclein Val110 variant gene.

XX Gamma-synuclein; human; single nucleotide polymorphism; SNP;

XX schizophrentia; neuroleptic; mutant; gene; ss.

XX Homo sapiens.

XX Key Location/Qualifiers
CDS 12..395

FT /*tag= a
FT /product= "Gamma-synuclein"
FT /transl_except= (pos:213..215,aa:Glu)
FT replace(340,A)
FT /*tag= b
FT /standard_name= "Single nucleotide polymorphism"

XX WO200275317-A2.

XX 26-SEP-2002.

XX 14-MAR-2002; 2002WO-EP002872.

XX 15-MAR-2001; 2001US-0276306P.

XX (NOVS) NOVARTIS AG.

XX (NOVS) NOVARTIS-ERFINDUNGEN VERW GES MBH.

XX (UYMA-) UNIV MARYLAND BALTIMORE.

XX Roberts RC, Van Oostrum J, Voshol J, Tamminga CA;

XX WPI; 2002-750574/81.

XX P-PsDB; ABP54932.

XX Screening for compounds for treating or interfering with the onset of
XX Schizophrenia Spectrum Disorders, by detecting interactions of candidate
XX compounds with the gamma-synuclein polypeptide.

XX Disclosure; Page; 32pp; English.

XX The present sequence is that of cDNA encoding the Val-110 isoform of
XX human gamma-synuclein. The invention relates to an isoform of gamma-
XX synuclein that is caused by an A/T single nucleotide polymorphism (SNP)
XX at position 329 of the gamma-synuclein coding sequence. This SNP causes a
XX glutamic acid to valine change at amino acid position 110 of gamma-

CC synuclein, and is associated with an increased susceptibility of
CC individuals to schizophrenia spectrum disorders (SSDs). This is the first
CC time that a genetic component of SSDs has been identified, and provides a
CC potential target for diagnosis and treatment of schizophrenia. Gamma-
CC synuclein polypeptides, especially those containing the E110V mutation,
CC are used in a claimed method of screening for compounds useful for the
CC treatment of SSDs, and gamma-synuclein expressing cells are used in a
CC claimed method of screening for agonist or antagonist compounds. An
CC oligonucleotide complementary to part of the gamma-synuclein coding
CC sequence is used for the discrimination of an SNP at position 329 of the
CC coding sequence. Gamma-synuclein polypeptides or polynucleotides are also
CC useful for the diagnosis of SSDs, or susceptibility to SSDs, e.g. by PCR
CC amplification of a polynucleotide encoding gamma-synuclein and analysis
CC of the occurrence of the SNP at position 329. A transgenic animal useful
CC for the study of SSDs is also claimed. Note: The present sequence is not
CC shown in the specification but is derived from the gamma-synuclein
CC sequence given in Fig 1 (see ABV73813)

CC
XX Sequence 550 BP; 131 A; 145 C; 192 G; 82 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.:	55.4	Length:	550
Score:	41.00	Matches:	8
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	6	Gaps:	0

US-09-017-715A-2_COPY_120_127 (1-8) x ABV73915 (1-550)

QY 1 GIUGUAAGInSerGIyASP 8
DB 369 GAGGAGGCCACAGTGGGGAGAC 392

RESULT 9
AAD63568
ID AAD63568 standard; cDNA; 550 BP.
AC AAD63568;
XX
XX AAD63568;
DT 12-FEB-2004 (first entry)
XX
XX Human amyloid-like protein cDNA.
DE
XX
XX Human; genetic disease; muscular dystrophy; cystic fibrosis; cytostatic;
KM scientific research; gene therapy; gene; amyloid-like protein; ss.
XX
XX Homo sapiens.
OS
XX
XX
FH Key Location/Qualifiers
FT CDS 12..395
FT /tag= a
FT /product= "Human amyloid-like protein"

XX
XX US6639052-B1.
XX
XX 28-OCT-2003.
XX
XX 14-OCT-1999; 99US-00417540.
XX
XX
XX 30-AUG-1995; 95US-0002993P.
XX 30-AUG-1996; 96US-00705771.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Moore PA;
XX
XX
XX WPI; 2003-842790/78.
XX P-PSDB; ABM02024.
XX
XX New isolated protein and nucleic acid molecules, useful for diagnostic
XX and therapeutic purposes, e.g. for treating genetic diseases such as
XX muscular dystrophy or cystic fibrosis.

XX
PS Example 5; Fig 1; Opp; English.
XX
XX The invention relates to isolated new isolated protein and nucleic acid
CC molecules useful for diagnostic and therapeutic purposes. The invention
CC is for treating genetic diseases such as muscular dystrophy or cystic
CC fibrosis, and for in vitro purposes related to scientific research,
CC synthesis of DNA and manufacture of DNA vectors. The invention is useful
CC in gene therapy. The present sequence is human amyloid-like protein cDNA

XX
XX Sequence 550 BP; 132 A; 145 C; 192 G; 81 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.:	55.4	Length:	550
Score:	41.00	Matches:	8
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	10	Gaps:	0

US-09-017-715A-2_COPY_120_127 (1-8) x AAD63568 (1-550)

QY 1 GIUGUAAGInSerGIyASP 8
DB 369 GAGGAGGCCACAGTGGGGAGAC 392

RESULT 10
ADG47636
ID ADG47636 standard; cDNA; 550 BP.
XX
XX ADG47636;
AC
XX
XX 11-MAR-2004 (first entry)
DT
XX
XX Human amyloid like protein cDNA.
DE
XX
XX ss; gene; muscular dystrophy; cystic fibrosis; hypertension;
KM angina pectoris; myocardial infarction; ulcer; asthma; allergy;
XX psychosis; depression; migraine; vomiting; benign prostatic hypertrophy;
XX osteoporosis; human.
XX
XX Homo sapiens.
OS
XX
XX
FH Key Location/Qualifiers
FT CDS 12..395
FT /tag= a
FT /product= "Amyloid like protein"

XX
XX US2003208043-A1.
XX
XX 06-NOV-2003.
XX
XX 04-JUN-2003; 2003US-00453478.
XX
XX 30-AUG-1995; 95US-0002993P.
XX 30-AUG-1996; 96US-00705771.
XX 14-OCT-1999; 99US-00417540.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Moore PA, Gentz RL, Ji H, Ni J, Hu J;
XX
XX WPI; 2003-864796/80.
XX P-PSDB; ADG47647.
XX
XX New human polypeptides and polynucleotides, useful for diagnosing or
XX treating genetic diseases such as muscular dystrophy or cystic fibrosis,
XX hypertension, asthma, depression or osteoporosis.
XX
XX Claim 18; SEQ ID NO 1; 56pp; English.
XX
XX The invention relates to an isolated human polypeptide. The polypeptides,
CC polynucleotides, agonists or antagonist are useful for diagnosing or

CC treating genetic diseases such as muscular dystrophy or cystic fibrosis,
CC hypertension, angina pectoris, myocardial infarction, ulcers, asthma,
CC allergies, psychoses, depression, migraine, vomiting, benign prostatic
CC hypertrophy or osteoporosis. The polypeptides and polynucleotides are
CC useful for in vitro purposes related to scientific research, synthesis of
CC DNA and manufacture of DNA vector. The present sequence represents cDNA
CC encoding human amyloid like protein.

SO Sequence 550 BP, 132 A, 145 C, 192 G, 81 T, 0 U, 0 Other;

Alignment Scores:

Pred. No.:	55.4	Length:	550
Score:	41.00	Matches:	8
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	10	Gaps:	0

US-09-017-715A-2_COPY_120_127 (1-8) x ADG47636 (1-550)

Qy 1 GUGUUAAGInserGlyGlyASP 8

Db 369 GAGGAGGCCCGAGACTGGGGGAGAC 392

RESULT 11

AAK29997
ID AAK29997 standard; DNA, 720 BP.

AC AAK29997;

DT 06-JUL-1999 (first entry)

DE Human persyn gene.

KM Human; synuclein; persyn; diagnosis; neurodegenerative disorder; cancer;

KM breast; skin; intermediate filament damage; ss.

OS Homo sapiens.

PN BP908727-A1.

PD 14-APR-1999.

PF 21-SEP-1998; 98BP-00307628.

PR 19-SEP-1997; 97GB-00019879.

PA (NEUR-) NEUROPA LTD.

(UYSA-) UNIV ST ANDREWS.

DR WPI; 1999-217169/19.

DR P-PSDB; AAY07271.

PT New synuclein protein (persyn) and gene, useful in assays for screening,

PT diagnosing or monitoring cancer, neurodegenerative disorders or skin

PT disorders.

PS Claim 29; Page 16-17; 39pp; English.

XX This sequence represents a novel human synuclein family

CC member designated persyn. The sequence is useful for screening,

CC diagnosing or monitoring cancer (especially breast or skin cancer),

CC neurodegenerative disorders or skin disorders and for identifying cells

CC having intermediate filament damage

XX Sequence 720 BP, 173 A, 209 C, 215 G, 123 T, 0 U, 0 Other;

Alignment Scores:

Pred. No.:	73.5	Length:	720
Score:	41.00	Matches:	8
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0

DB: 2 Gaps: 0

US-09-017-715A-2_COPY_120_127 (1-8) x AAK29997 (1-720)

Qy 1 GUGUUAAGInserGlyGlyASP 8

Db 406 GAGGAGGCCCGAGACTGGGGGAGAC 429

RESULT 12

ABK76519
ID ABK76519 standard; cDNA, 720 BP.

AC ABK76519;

DT 11-DEC-2002 (first entry)

DE cDNA encoding human ovarian cancer marker OV60.

KM Human; ovarian cancer; marker; cancer; familial history; brain disorder;

KM central nervous system disorder; bacterial meningitis; viral meningitis;

KM Alzheimer's disease; Parkinson's disease; cerebral oedema; hydrocephalus;

KM brain herniation; inflammation; encephalitis; testicular disorder;

KM non-tuberculous granulomatous orchitis; connective tissue disorder;

KM heart disorder; ischaemic heart disease; atherosclerosis; neoplasm;

KM histological type; carcinogenic; ovarian cancer marker; gene; ss.

OS Homo sapiens.

PN WO200271928-A2.

PD 19-SEP-2002.

PF 14-MAR-2002; 2002MO-US007826.

PR 14-MAR-2001; 2001US-0276025P.

PR 14-MAR-2001; 2001US-0276026P.

PR 10-AUG-2001; 2001US-0311732P.

PR 19-SEP-2001; 2001US-0323580P.

PR 26-SEP-2001; 2001US-0324967P.

PR 26-SEP-2001; 2001US-0325102P.

PR 26-SEP-2001; 2001US-0325149P.

XX (MIL-) MILBENTUM PHARM INC.

PI Monahan JE, Gannavarapu M, Hoersch S, Kamatkar S, Kovatis SG;

PI Meyers RE, Morrissey MP, Olandt PJ, Sen A, Vieby PO, Mills GB;

PI Bast RC, Lu K, Schmandt RB, Zhao X, Glatz K;

DR WPI; 2002-723277/78.

DR P-PSDB; ABG96420.

PT Assessing whether a patient is afflicted with ovarian cancer, useful in

PT assessing the stage or progression of the disease, compares comparing

PT the expression level of a cancer marker in a sample from a patient and

PT from a non cancer patient.

PS Disclosure; Page 411; 481pp; English.

XX The present invention relates to a new method for assessing whether a

CC patient is afflicted with ovarian cancer. The method involves comparing

CC the expression level of a marker in a patient sample and the normal level

CC of expression of the marker in a control non-ovarian cancer sample, where

CC the marker is selected from 363 cancer markers described in the

CC specification. The method of the invention is useful in diagnosing or

CC characterising cancer, in detecting the presence of cancer as early as

CC possible, and the recurrence of ovarian cancer. The method may also be of

CC particular use with patients having an enhanced risk of developing

CC ovarian cancer (e.g. patients having a familial history of ovarian

CC cancer). The cancer markers may be used in the management and treatment

CC of e.g. brain and central nervous system disorders (e.g. bacterial and

CC viral meningitis, Alzheimer's disease or Parkinson's disease), brain

CC disorders (e.g. cerebral oedema, hydrocephalus or brain herniations),

CC inflammations (e.g. bacterial or viral meningitis or encephalitis),

XX Claim 84; SEQ ID NO 469; 205bp; English.
PS
XX This invention relates to a novel method of determining a predisposition
CC for or the occurrence of neurodegenerative disease comprising detecting
CC in a target nucleic acid obtained from the subject the presence of an
CC allelic variant of polymorphic regions of human genes selected from
CC uridine kinase plasmidogen activator (uPA), gamma-globulin (SNGC), insulin
CC degrading enzyme (IDE), kinesin-like protein 1 (KSL1), lysosomal acid
CC lipase (LIPA) and tumour necrosis factor receptor SF6 (TNFRSF6). The
CC method is useful in determining the presence or predisposition to a
CC neurodegenerative disease, particularly Alzheimer's disease. The present
CC sequence is the cDNA sequence of the human SNGC gene which is related to
CC the invention.
XX
SQ Sequence 720 BP, 172 A; 205 C; 212 G; 120 T; 0 U; 11 Other;
Alignment Scores:
Pred. No.: 73.5 Length: 720
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 12 Gaps: 0
US-09-017-715A-2_COPY_120_127 (1-8) x ADHS4342 (1-720)
QY 1 GluGluAlaGlnSerGlyIAsp 8
Db 406 GAGGAGGCCCGAGTGGGGGAGAC 429
RESULT 15
AAI93778
ID AAI93778 standard; cDNA; 783 BP.
XX
XX AAI93778;
AC
XX
XX 06-NOV-2001 (first entry)
DT
XX
XX Human polynucleotide SEQ ID NO 13838.
DE
XX
XX Human; cytokine; cell proliferation; cell differentiation; gene therapy;
KM vaccine; peptide therapy; stem cell growth factor; haematopoiesis;
KM tissue growth factor; immunomodulatory; cancer; leukemia;
KM nervous system disorders; arthritis; inflammation; ss.
XX
XX Homo sapiens.
OS
XX
XX WO200164835-A2.
PN
XX
XX 07-SEP-2001.
PD
XX
XX 26-FEB-2001; 2001WO-US004927.
PF
XX
XX 28-FEB-2000; 2000US-00515126.
PR
XX 18-MAY-2000; 2000US-00577409.
PA
XX (HYSE-) HYSEQ INC.
XX
XX Tang YT, Liu C, Drmanac RT;
PI
XX WPI: 2001-514838/56.
DR P-PSDB; AAO13847.
XX
XX Isolated nucleic acids and polypeptides, useful for preventing diagnosing
PT and treating e.g. leukemia, inflammation and immune disorders.
XX
XX Claim 1; SEQ ID NO 13838; 1399bp + Sequence Listing; English.
PS
XX The invention relates to human polynucleotides (AAI79941-AAI93841) and
CC the encoded proteins (AAO00010-AAO013910) that exhibit activity elating to
CC cytokine, cell proliferation or cell differentiation or which may induce
CC production of other cytokines in other cell populations. The

CC polynucleotides and polypeptides are useful in gene therapy, vaccines or
CC peptide therapy. The polypeptides have various cytokine-like activities,
CC e.g. stem cell growth factor activity, haematopoiesis regulating
CC activity, tissue growth factor activity, immunomodulatory activity and
CC activin/inhibin activity and may be useful in the diagnosis and/or
CC treatment of cancer, leukemia, nervous system disorders, arthritis and
CC inflammation. Note: the sequence data for this patent did not form part
CC of the printed specification, but was obtained in electronic format
CC directly from WIPO at ftp.wipo.int/pub/published_pct_sequences
XX
SQ Sequence 783 BP, 187 A; 232 C; 237 G; 127 T; 0 U; 0 Other;
Alignment Scores:
Pred. No.: 80.3 Length: 783
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0
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Db 471 GAGGAGGCCCGAGTGGGGGAGAC 494
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(without alignments)
706.682 Million cell updates/sec

Title: US-09-017-715A-2

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-FCAGPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	610	100.0	550	4	US-09-417-540-1
3	602	98.7	702	4	US-09-949-016-1915
4	595	97.5	720	4	US-09-949-016-442
5	468.5	76.8	786	5	PCT-US95-08295-1
6	307.5	50.4	1543	4	US-09-949-016-58
7	307.5	50.4	1543	4	US-09-949-016-3084
8	307.5	50.4	1543	4	US-09-949-016-3085
9	307.5	50.4	1560	5	PCT-US94-09789-1
10	303.5	49.8	1096	4	US-09-949-016-3086
11	303.5	49.8	1096	4	US-09-949-016-3087
12	284.5	46.6	703	4	US-09-949-016-2926

13	282	46.2	313	4	US-09-513-999C-2232	Sequence 2232, Ap
14	201.5	33.0	8607	4	US-09-949-016-13657	Sequence 13657, A
15	201.5	33.0	8608	4	US-09-949-016-12184	Sequence 12184, A
16	187	30.7	419	4	US-09-621-976-609	Sequence 609, App
17	181	30.2	409	4	US-09-513-999C-2347	Sequence 2347, Ap
18	171	28.0	8281	4	US-09-949-016-14668	Sequence 14668, A
19	155	25.4	113876	4	US-09-949-016-14828	Sequence 14828, A
20	155	25.4	113876	4	US-09-949-016-14829	Sequence 14829, A
21	155	25.4	115508	4	US-09-949-016-11800	Sequence 11800, A
22	155	25.4	115508	4	US-09-949-016-14826	Sequence 14826, A
23	155	25.4	115508	4	US-09-949-016-14827	Sequence 14827, A
24	113	18.5	601	4	US-09-949-016-18715	Sequence 18715, A
25	113	18.5	601	4	US-09-949-016-111738	Sequence 111738, A
26	113	18.5	601	4	US-09-949-016-111884	Sequence 111884, A
27	113	18.5	601	4	US-09-949-016-111884	Sequence 111884, A
28	113	18.5	601	4	US-09-949-016-111884	Sequence 111884, A
29	98	16.1	845	5	PCT-US96-05320A-1342	Sequence 1342, Ap
30	98	16.1	845	5	PCT-US96-05320A-1342	Sequence 25, Appl
31	98	16.1	845	3	US-08-743-6378-25	Sequence 25, Appl
32	98	16.1	1830121	4	US-09-557-884-1	Sequence 1, Appl
33	98	16.1	1830121	4	US-09-557-884-1	Sequence 1, Appl
34	98	16.1	1830121	4	US-09-643-990A-1	Sequence 1, Appl
35	98	16.1	1830121	4	US-09-643-990A-1	Sequence 1, Appl
36	95	15.6	729	4	US-09-543-681A-1761	Sequence 1761, Ap
37	94	15.4	1695	2	US-08-216-894-1	Sequence 1, Appl
38	94	15.4	1695	2	US-09-115-746-1	Sequence 7, Appl
39	94	15.4	1932	2	US-08-216-894-7	Sequence 7, Appl
40	94	15.4	1932	3	US-09-115-746-7	Sequence 2, Appl
41	94	15.4	5361	3	US-08-973-462-2	Sequence 1, Appl
42	94	15.4	6152	1	US-08-973-462-1	Sequence 1903, Ap
43	93	15.2	6717	4	US-09-107-433-1903	Sequence 1385, Ap
44	92.5	15.2	2787	4	US-09-134-000C-3185	Sequence 3423, Ap
45	92	15.1	3506	4	US-09-710-279-3423	

ALIGNMENTS

RESULT 1
US-08-705-771-1
Sequence 1, Application US/08705771
Patent No. 6054289
GENERAL INFORMATION:
APPLICANT: Paul Moore, Reiner Gentz, Hongjin Ji,
TITLE OF INVENTION: Human Genes, Sequences and
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: CECCHI, BYRNE, BAIN, GILFILLAN,
STREET: 6 BECKER FARM ROAD
CITY: ROSELAND
STATE: NEW JERSEY
COUNTRY: USA
ZIP: 07068
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5 INCH DISKETTE
COMPUTER: IBM PS/2
OPERATING SYSTEM: MS-DOS
SOFTWARE: WORD PERFECT 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/705,771
FILING DATE: August 30, 1996
CLASSIFICATION: 536
ATTORNEY/AGENT INFORMATION:
NAME: MULHINS, J.G.
REGISTRATION NUMBER: 33,073
REFERENCE/DOCKET NUMBER: 325800-346 (PFI96)
TELECOMMUNICATION INFORMATION:
TELEPHONE: 973-994-1744
TELEFAX: 973-994-1744
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:

LENGTH: 550 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-705-771-1

Alignment Scores:

Pred. No.:	2,156-69	Length:	550
Score:	610.00	Matches:	127
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	3	Gaps:	0

US-09-017-715A-2 (1-127) x US-08-705-771-1 (1-550)

QY 1 Metaspvalpshelyslypheserilealylysglyvalvalglvalglu 20
DB 12 ATGAGTGTTCAGAGAGGGCTTCTCCATCGCCAGAGAGGGGTGGTGGTGGAA 71
QY 21 LysThrLysGlnGlyValThrGluAlaGluLysThrLysGlnGlyValMetYrVal 40
DB 72 AAGACCAAGCAGGGGTGACGAGACGCTGAGAACCAAGAGGGGTGATGTGTG 131
QY 41 G1yAlaLysThrLysGluAsnValGlnSerValThrSerValAlaGluLysThrLys 60
DB 132 GAGGCCAAGACCAAGAGAGATGTTGTACAGCGCTCACTGAGCCCAAGACCAAG 191
QY 61 G1uGlnAlaAsnAlaValSerLysAlaValSerSerValAsnThrValAlaThrLys 80
DB 192 GAGCAGGCCAAGCGCTGAGCAAGGCTGTGTGACAGCGTCAACACTGTGGCCACCAAG 251
QY 81 ThrValGlnGluAlaGluAsnIleAlaValThrSerGlyValAlaArgLysGluAspLeu 100
DB 252 ACCGTGAGAGAGCGAGAACATCGCGTCACTCGGGGTGTGGCAAGAGAGACTTG 311
QY 101 ArgProSerAlaProGlnGlnGluAlaSerLysGluLysGlnGluValAlaGlu 120
DB 312 AGGCCATCTGCCCCCAACAGAGGGGTGAGCATCCAAAGAGAAAGAGAGTGGCAGAG 371
QY 121 G1uAlaGlnSerGlyLysp 127
DB 372 GAGGCCAAGATGGGGAGAGAC 392

RESULT 2

US-09-417-540-1
Sequence 1, Application US/09417540
Patent No. 6639052

GENERAL INFORMATION:

APPLICANT: Paul Moore, Reiner Gentz, Hongjin Ji,
Jian Ni and Jing-Shan Hu
TITLE OF INVENTION: Human Genes, Sequences and
Expression Products
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: CERCHI, STEWART & OLSTEIN,

STREET: 6 BECKER FARM ROAD
CITY: ROSELAND
STATE: NEW JERSEY
COUNTRY: USA
ZIP: 07068

COMPUTER READABLE FORM:

MEDIUM TYPE: 3.5 INCH DISKETTE
COMPUTER: IBM PS/2
OPERATING SYSTEM: MS-DOS
SOFTWARE: WORD PERFECT 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/417,540
FILING DATE: 14-Oct-1999
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/705,771
FILING DATE: August 30, 1996
ATTORNEY/AGENT INFORMATION:
NAME: MULLINS, J.G.

REGISTRATION NUMBER: 33,073
REFERENCE/DOCKET NUMBER: 325800-346 (PFI96)
TELECOMMUNICATION INFORMATION:
TELEPHONE: 973-994-1700
TELEFAX: 973-994-1744

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:
LENGTH: 550 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear

MOLECULE TYPE: DNA

SEQUENCE DESCRIPTION: SEQ ID NO: 1:

US-09-417-540-1

Alignment Scores:
Pred. No.: 2,156-69 Length: 550
Score: 610.00 Matches: 127
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-09-017-715A-2 (1-127) x US-09-417-540-1 (1-550)

QY 1 Metaspvalpshelyslypheserilealylysglyvalvalglvalglu 20
DB 12 ATGAGTGTTCAGAGAGGGCTTCTCCATCGCCAGAGAGGGGTGGTGGTGGAA 71
QY 21 LysThrLysGlnGlyValThrGluAlaGluLysThrLysGlnGlyValMetYrVal 40
DB 72 AAGACCAAGCAGGGGTGACGAGACGCTGAGAACCAAGAGGGGTGATGTGTG 131
QY 41 G1yAlaLysThrLysGluAsnValGlnSerValThrSerValAlaGluLysThrLys 60
DB 132 GAGGCCAAGACCAAGAGAGATGTTGTACAGCGTCACTGAGCCCAAGACCAAG 191
QY 61 G1uGlnAlaAsnAlaValSerLysAlaValSerSerValAsnThrValAlaThrLys 80
DB 192 GAGCAGGCCAAGCGCTGAGCAAGGCTGTGTGACAGCGTCAACACTGTGGCCACCAAG 251
QY 81 ThrValGlnGluAlaGluAsnIleAlaValThrSerGlyValAlaArgLysGluAspLeu 100
DB 252 ACCGTGAGAGAGCGAGAACATCGCGTCACTCGGGGTGTGGCAAGAGAGACTTG 311
QY 101 ArgProSerAlaProGlnGlnGluAlaSerLysGluLysGlnGluValAlaGlu 120
DB 312 AGGCCATCTGCCCCCAACAGAGGGGTGAGCATCCAAAGAGAAAGAGAGTGGCAGAG 371
QY 121 G1uAlaGlnSerGlyLysp 127
DB 372 GAGGCCAAGATGGGGAGAGAC 392

RESULT 3

US-09-949-016-1915
Sequence 1915, Application US/09949016
Patent No. 6812339

GENERAL INFORMATION:

APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
FILE REFERENCE: CL001307
CURRENT APPLICATION NUMBER: US/09/949,016

PRIOR FILING DATE: 2000-04-14
CURRENT APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 60/231,498


```
DB 95 ATGGATGTTTCAGAGGGCTTCTCCATGCCAGAGAGGGCGTGGTGGCGGAGAA 154
QY 21 LYSTHLYSGINGLYVALTHRGUAAAGLULVSTHLYSGINGLYVALMETYRVAL 40
DB 155 AAGACCAACAGAGGGGTGACGAGAGCTGAGAAACCAAGAGGGGTCTATGTATGTG 214
QY 41 GYALALYSTHLYSGUASN--VAL-VALGINSERVALTHRSERVALAGLULYSTH 59
DB 215 GGAGCCAAACCAAGAGAGATGTTGATATGACAGACCGTCACTGCTGCCGAGAGAC 274
QY 59 RLYSGUAGUAAASNAVALSERLYSALVALSERSERVALASRTHVALALATH 79
DB 275 CAAGGCGAGGCGCAACGCGGTGAGCAAGCGTGTGTGACAGCGTCAACACTKGTGCCAC 334
QY 79 RLYSTHVALGUGUAGUAGUASNLVALVALTHRSERGLYVALARGLYSGUAS 99
DB 335 CAAGACCGTGAAGAGGCGAGAACATCGCGGTCAACCTCCGGGKTGCTCGCAAGAGGA 394
QY 99 PLEUATGPROSERALAPROGLINGLUGLYGUALASERLYSGLU-LYSGUGUVALA 119
DB 395 YTTKAGCCCATY-TKCCCCCAACAGAGGGTGAAGCATMAAAGAAARAGAKXGSAAGWG 453
QY 119 IAGUGUAGUAGINSERGLYLYASP 127
DB 454 CMRAKRGMSCAAGAGTGGGGGAGAC 479
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RESULT 6

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US-09-949-016-58
/ Sequence 58, Application US/09949016
/ Patent No. 6812339
/ GENERAL INFORMATION:
/ APPLICANT: VENTER, J. Craig et al.
/ TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
/ TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
/ FILE REFERENCE: CL001307
/ CURRENT APPLICATION NUMBER: US/09/949,016
/ PRIOR FILING DATE: 2000-04-14
/ PRIOR APPLICATION NUMBER: 60/241,755
/ PRIOR FILING DATE: 2000-10-20
/ PRIOR APPLICATION NUMBER: 60/237,768
/ PRIOR FILING DATE: 2000-10-03
/ PRIOR APPLICATION NUMBER: 60/231,498
/ PRIOR FILING DATE: 2000-09-08
/ NUMBER OF SEQ ID NOS: 207012
/ SOFTWARE: FastSeq for Windows Version 4.0
/ SEQ ID NO: 58
/ LENGTH: 1543
/ TYPE: DNA
/ ORGANISM: Human
US-09-949-016-58
```

Alignment Scores:

Pred. No.:	8.5e-30	Length:	1543
Score:	307.50	Matches:	68
Percent Similarity:	70.91%	Conservative:	10
Best Local Similarity:	61.82%	Mismatches:	29
Query Match:	50.41%	Indels:	3
DB:	4	Gaps:	1

US-09-017-715A-2 (1-127) x US-09-949-016-58 (1-1543)

```
QY 1 MetaspvalPheLysGlyPheSerIleAlaLysLysGlyValValGlyValAlaGlu 20
DB 47 ATGGATGATTCATGAAAGCACTTTAAAGCCAAAGAGAGAGTGTGCTGCTGTGAG 106
QY 21 LYSTHLYSGINGLYVALTHRGUAAAGLULVSTHLYSGINGLYVALMETYRVAL 40
DB 107 AAAACCAACAGAGGGTGTGACAGAGCAGAGAAACAAAGAGGGTCTTCTATGTA 166
QY 41 GYALALYSTHLYSGUASNVALGINSERVALTHRSERVALAGLULYSTHLYS 60
DB 167 GGCTCCAAAACCAAGAGAGAGTGTGATGTGTGCAACAGTGTGAGAAAGACCAA 226
```

```
QY 61 GUUGUAAASNAVALSERLYSALVALSERSERVALASRTHVALALATHLYS 80
DB 227 GAGCAAGAGCAAAAGTTGAGAGACAGTGTGAGAGGGGTGACAGCACTAGCCAGAA 286
QY 81 THVALGUGUAGUAGUASNLVALVALTHRSERGLYVALARGLYSGUASPLEU 100
DB 287 ACAGTGAAGGAGGAGGAGCATTTGCACAGCCACTGCTTGTCTCAAAAAGACCAATTG 346
QY 101 -----ArgProSerAlaProGlnGln 107
DB 347 GGCAAGAAATGAAGAGAGCCCAAGGAA 376
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RESULT 7

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US-09-949-016-3084
/ Sequence 3084, Application US/09949016
/ Patent No. 6812339
/ GENERAL INFORMATION:
/ APPLICANT: VENTER, J. Craig et al.
/ TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
/ TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
/ FILE REFERENCE: CL001307
/ CURRENT APPLICATION NUMBER: US/09/949,016
/ PRIOR FILING DATE: 2000-04-14
/ PRIOR APPLICATION NUMBER: 60/241,755
/ PRIOR FILING DATE: 2000-10-20
/ PRIOR APPLICATION NUMBER: 60/237,768
/ PRIOR FILING DATE: 2000-10-03
/ PRIOR APPLICATION NUMBER: 60/231,498
/ PRIOR FILING DATE: 2000-09-08
/ NUMBER OF SEQ ID NOS: 207012
/ SOFTWARE: FastSeq for Windows Version 4.0
/ SEQ ID NO: 3084
/ LENGTH: 1543
/ TYPE: DNA
/ ORGANISM: Human
US-09-949-016-3084
```

Alignment Scores:

Pred. No.:	8.5e-30	Length:	1543
Score:	307.50	Matches:	68
Percent Similarity:	70.91%	Conservative:	10
Best Local Similarity:	61.82%	Mismatches:	29
Query Match:	50.41%	Indels:	3
DB:	4	Gaps:	1

US-09-017-715A-2 (1-127) x US-09-949-016-3084 (1-1543)

```
QY 1 MetaspvalPheLysGlyPheSerIleAlaLysLysGlyValValGlyValAlaGlu 20
DB 47 ATGGATGATTCATGAAAGCACTTTAAAGCCAAAGAGAGAGTGTGCTGCTGTGAG 106
QY 21 LYSTHLYSGINGLYVALTHRGUAAAGLULVSTHLYSGINGLYVALMETYRVAL 40
DB 107 AAAACCAACAGAGGGTGTGACAGAGCAGAGAAACAAAGAGGGTCTTCTATGTA 166
QY 41 GYALALYSTHLYSGUASNVALGINSERVALTHRSERVALAGLULYSTHLYS 60
DB 167 GGCTCCAAAACCAAGAGAGAGTGTGATGTGTGCAACAGTGTGAGAAAGACCAA 226
QY 61 GUUGUAAASNAVALSERLYSALVALSERSERVALASRTHVALALATHLYS 80
DB 227 GAGCAAGAGCAAAAGTTGAGAGACAGTGTGAGAGGGGTGACAGCACTAGCCAGAA 286
QY 81 THVALGUGUAGUAGUASNLVALVALTHRSERGLYVALARGLYSGUASPLEU 100
DB 287 ACAGTGAAGGAGGAGGAGCATTTGCACAGCCACTGCTTGTCTCAAAAAGACCAATTG 346
QY 101 -----ArgProSerAlaProGlnGln 107
DB 347 GGCAAGAAATGAAGAGAGCCCAAGGAA 376
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RESULT 8

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US-09-949-016-3085
; Sequence 3085, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949, 016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FASTSEQ for Windows Version 4.0
; SEQ ID NO 3085
; LENGTH: 1543
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-3085

Alignment Scores:
Pred. No.: 8,5e-30 Length: 1543
Score: 307.50 Matches: 68
Percent Similarity: 70.91% Conservative: 10
Best Local Similarity: 61.82% Mismatches: 29
Query Match: 50.41% Indels: 3
DB: 4 Gaps: 1

US-09-017-715A-2 (1-127) x US-09-949-016-3085 (1-1543)
QY 1 MetAspValPheIySgIyPheSerIleAlaIySgIyValIglIyAlaIglu 20
DB 47 ATGGATGATTCATGAAGAAGCTTCAAGGCCAAGGAGGAGTGTGGCTGCTGAG 106
QY 21 LyeThrIySgIyValIthrGluAlaIgluIyThrIySgIyValIgluIyMetTyVal 40
DB 107 AAAACCAAGGCTGTGGCAGAGCAGAAAGCAAAAGGCTGTCTATGTA 166
QY 41 G1yAlaIyThrIySgIyValIthrGluAlaIgluIyThrIySgIyValIgluIyThrIy 60
DB 167 GGCTCCAAACCAAGGAGGAGTGTGATGCTGTGCAACAGTGGCTGAGAAACCAA 226
QY 61 GluGlnAlaAsnAlaIySerIyAlaIyAlaIySerIyAlaIyThrIySgIyValIthrIyS 80
DB 227 GAGCAAGTGAACAATGTTGAGAGCAGTGTGACGGGTGACAGCAGTACCCAGAG 286
QY 81 ThrValIgluIyAlaIyAsnIleAlaIyValIthrSergIyValIyAlaIyIySgIyAlaIy 100
DB 287 ACAGTGAGGAGGAGGAGGAGTGTGACAGCCACGCTTGTTCAAAAGAACCACTTG 346
QY 101 -----ArgProSerAlaProGlnI 107
DB 347 GGCAGAGATGAAGAGAGCCCAAGGAA 376

RESULT 9
PCT-US94-09789-1
; Sequence 1, Application PC/TUS9409789
; GENERAL INFORMATION:
; APPLICANT: The Regents of the University of California
; TITLE OF INVENTION: NOVEL COMPONENT OF AMYLOID IN
; TITLE OF INVENTION: ALZHEIMER'S DISEASE AND METHODS FOR USE OF SAME
; NUMBER OF SEQUENCES: 12
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Spensley Horn Jubas & Lubitz
; STREET: 1880 Century Park East - Suite 500
; CITY: Los Angeles
; STATE: California
; COUNTRY: USA
; ZIP: 90067
```

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COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US94/09789
; FILING DATE: 29-AUG-1994
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Howells, Stacy L.
; REGISTRATION NUMBER: 34,842
; REFERENCE/DOCKET NUMBER: PD-3520
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 455-5100
; TELEFAX: (619) 455-5110
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1560 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; IMMEDIATE SOURCE:
; CLONE: cDNA for NACP
; FEATURE:
; NAME/KEY: misc RNA
; LOCATION: 1..1560
PCT-US94-09789-1

Alignment Scores:
Pred. No.: 8,64e-30 Length: 1560
Score: 307.50 Matches: 68
Percent Similarity: 70.91% Conservative: 10
Best Local Similarity: 61.82% Mismatches: 29
Query Match: 50.41% Indels: 3
DB: 5 Gaps: 1

US-09-017-715A-2 (1-127) x PCT-US94-09789-1 (1-1560)
QY 1 MetAspValPheIySgIyPheSerIleAlaIySgIyValIglIyAlaIglu 20
DB 53 ATGGATGATTCATGAAGAAGCTTCAAGGCCAAGGAGGAGTGTGGCTGCTGAG 112
QY 21 LyeThrIySgIyValIthrGluAlaIgluIyThrIySgIyValIgluIyMetTyVal 40
DB 113 AAAACCAAGGCTGTGGCAGAGCAGAAAGCAAAAGGCTGTCTATGTA 172
QY 41 G1yAlaIyThrIySgIyValIthrGluAlaIgluIyThrIySgIyValIgluIyThrIy 60
DB 173 GGCTCCAAACCAAGGAGGAGTGTGATGCTGTGCAACAGTGGCTGAGAAACCAA 232
QY 61 GluGlnAlaAsnAlaIySerIyAlaIyAlaIySerIyAlaIyThrIySgIyValIthrIyS 80
DB 233 GAGCAAGTGAACAATGTTGAGAGCAGTGTGACGGGTGACAGCAGTACCCAGAG 292
QY 81 ThrValIgluIyAlaIyAsnIleAlaIyValIthrSergIyValIyAlaIyIySgIyAlaIy 100
DB 293 ACAGTGAGGAGGAGGAGGAGTGTGACAGCCACGCTTGTTCAAAAGAACCACTTG 352
QY 101 -----ArgProSerAlaProGlnI 107
DB 353 GGCAGAGATGAAGAGAGCCCAAGGAA 382

RESULT 10
US-09-949-016-3086
; Sequence 3086, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
```


Db	418	ATGAGCGTGTTCATGAAAGGGCCCTGTCCATATGGCCAAAGAGGGGGCTGTGTGGACGCCCGAG	477
Qy	21	lyeThrIyrgslnglnlyValThrGluAlaAlaGluIyThrIyGluGluIyAlaMecTyVal	40
Db	478	AAACCAAGACGAGGGGCTGCACCGAGGGGGGGAGAAACCAAGAGAGGGGCTCTTACGTC	533
Qy	41	GlyAlaIyIyThrysgIuBsnValValGlnSerValThrSerValAlGluIyThrIys	60
Db	538	GGAAGCAAGACCCGAGAAGGTGTGTACAAAGGTGTGGCTTCAGTGGCTGAAAAACCAAG	597
Qy	61	GluGlnAlaAsnAlaValSerIyValAlaValSerSerValAsnThrValAlaThrIys	80
Db	598	GAACAGGCCCTCCATCTCGAGAGAGCTGTGTTCTCTGGG-----	636
Qy	81	ThrValGluGluIyIyGluBsnIleAlaValThrSerGlyValIyValIyGluIyAsp	99
Db	637	-----GCAAGGAACATCGACGACCAACGAGCTGTGTAAAGAGGAGGA	681

RESULT 13
US-09-513-999C-2232
; Sequence 2232, Application US/09513999C
Patent No. 679001

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? PATENT NO.: 6783961
? GENERAL INFORMATION:
? APPLICANT: Dumas Milne Edwards, J.B.
? APPLICANT: Duclert, A.
? APPLICANT: Giordano, J.Y.
? TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins
? Patent No. 6783961
? FILE REFERENCE: 59.US2.REG
? CURRENT APPLICATION NUMBER: US/09/513,999C
? CURRENT FILING DATE: 2000-02-24
? PRIOR APPLICATION NUMBER: US 60/122,487
? PRIOR FILING DATE: 1999-02-26
? NUMBER OF SEQ ID NOS: 36681
? SOFTWARE: Patent.pm
? SEQ ID NO 2232
? LENGTH: 313
? TYPE: DNA
? ORGANISM: Homo sapiens
? FEATURE:
? NAME/KEY: CDS
? LOCATION: 136..312
? FEATURE:
? NAME/KEY: misc_feature
? LOCATION: 117..feature
? OTHER INFORMATION: m=a or c
? OS-09-513-999C-2232

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Alignment Scores:			
Pred. No.:	1,56e-27	length:	311
Score:	282.00	Matches:	58
Percent Similarity:	100.00%	Conservative:	1
Best local Similarity:	98.31%	Mismatches:	0
Query Match:	46.23%	Indels:	0
DB:	4	Gaps:	0

Oy	MeAspAlaPheIleYstIysGlyPheSerIleAlaIleValYsGIValValIGIValIValGIu	20
	
Db	ATGATGTCCTTCAAGAAAGGAGCTTCTCCATCGCCCAAGAGAGGCGTGTGGTGGCGGTGGAA	195
Oy	136	
	21	
Db	196	
Oy	41	
Db	256	

RESULT 14
US-09-949-016-13657
; Sequence 13657, Application US/09949016

```

; Patient No:6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: C1001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FASTSEQ for Windows Version 4.0
; SEQ ID NO 13657
; LENGTH: 8607
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)_(8607)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-13657

Alignment Scores:
Pred. No.:          4,69e-15          Length:      8607
Score:              201.50            Matches:      97
Percent Similarity: 20.66%            Conservative: 3
Best Local Similarity: 20.04%         Mismatches:  3
Query Match:        33.03%            Indels:      33
DB:                  4                Gaps:        2

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US-09-949-016-13657

Alignment Scores:

Pred. No.:	4,696-15	Length:	8607
Score:	201.50	Matches:	97
Percent Similarity:	20.66%	Conservative:	3
Best local Similarity:	20.04%	Mismatches:	34
Query Match:	33.03%	Indels:	383
DB:	4	Gaps:	2

US-09-017-715A-2 (1-127) x US-09-949-016-13657 (1-8607)

QY 1 MetAspValPheLysLysGlyPheSerIleAlaLysLysGlyValValGlyAlaValGlu 20

Db 2048 ATGGATGTCCTCAAGAAGGCTTCTCCATCGCCAAGGAGGCGGTGGTGGGTGCCGTGAA 2107

21 LysThrLysGlnGlyValThrGluAlaIaGluLysThrLysGlnGlyValMetCysVal 40

Db 2108 AAGACCAAGCAGGGGTGACGAGCAGCTGAGAAGACCAAGGAGGGGTCATGTATGTG 21

40 ----- 40

Db 2168 GGTAGTGGGCATGGCAGGTTGGACAGTGTGGTGGCCAAAGGTGAGTGCCCACTAC 2227

q_Y 40 ----- 40

Db 2228 CTTGCCAGACCTTACTCCCCAGCCCCAGGAGGCATTTTGGAGGGGGCGAGCCCTGG 22

q_Y 40 ----- 40

Db 2288 CTATCAGGTGGGTCTCCAGACCCCTGAGCACCCACAATGCCCTGTGCACACCTATGTG 23

40 ----- 40

Db 2348 TGTTCCTTGGCCTCTCGGCTCTGGGTGT CAGAGACCGCAGACAGGGCTGCTA 24

40 ----- 40

Db 2408 CCTGTCGTGCACGACACACATTCGCCAAGCATACCGCCTCCCTGAGCCTGAGCC 24

40 ----- 40

Db 2468 CCTGAAGCCATGAGCAGCCTGTGCTCAGGTGCCCCCAACCCTCTCCACACGGGAGCGCT 25

QY 40 ----- 40

Db 2528 ACAGCCAGTCA CGATCCCTCTCCCGAGAGAGAGGGCAGGCTGGGGATGAAC 25

40 ----- 40

```

Db      2588 CTAGGCTAGTGTTCCTCCCTCCCGCATCTCTCTGACACTCTCCAGAGAGAGAGGGA 2647
Qy      40 ----- 40
Db      2648 GGTCAAGCAATGACTCAGCTCTGGCCCATCTGTCCTGCTGCTCTGAGGCCGGC 2707
Qy      40 ----- 40
Db      2708 CACACCCGGGCAAGGGGCTGGACCTGGGGTCTAGCCAGTGTCTTACCTCAGGCTGTCT 2767
Qy      40 ----- 40
Db      2768 CTCTTGTCCCAACATTCCTGCTGCTGCTCCCTTCCATTCATCATTCTTCCAGACAG 2827
Qy      40 ----- 40
Db      2828 CAGGAAGGCGCTCTGAAGGGGCGCGCGCCCGACAGACATCCTTACCCCGCCACCG 2887
Qy      40 ----- 40
Db      2888 ACCCCACAGTTTGTCCAGCTGTTCTGTTGTTTGTCTGACCGCCCAACACCTCGAG 2947
Qy      41 ----- 49
Db      2948 GGAGGCTGGGCTGACAGCTCATTTCTCCCGAGGACCAAGACAGAGATGTTCT 3007
Qy      49 |G|n|s|e|r|v|a|l|t|h|y|s|e|r|l|e|a|l|y|s|g|l|y|v|a|l|g|l|u 54
Db      3008 ACAGAGCGTGAACCTC-AGGTGAGAAAGCCCGAGGCGGACACATGGGGATAGAGACC 3066
Qy      54 ----- 54
Db      3067 CCTGGGGCTCTGCACTCTAGTGTGGGGCTCAAACTAGAGTCTGCTTACCCCAAC 3126
Qy      54 ----- 54
Db      3127 TGGGGTCCAGAGCCCTACAGACCCCTGACAGACATGAGGCTAACTAGGTGGGGCTTC 3186
Qy      54 ----- 54
Db      3187 CTTACCCCAACGATCAGAGGTGCTGTGATCAGAGGAGGAGGGATGCCAGC 3246
Qy      54 ----- 54
Db      3247 AAGGCCAGGGCTCTGAGCTCTGGAGAGGGGCTGCGAGCTGACTCAGACAGGCTGCT 3306
Qy      55 ----- 60
Db      3307 TGGGGCTGGGGCTGGGGGTGAGGCGCAGCGATGCTCTCCCATAGTGGCCGAGAAACCAA 3366
Qy      60 |s|g|l|u|a|a|s|n|a|v|a|l|s|e|r|v|a|l|s|e|r|v|a|l|s|e|r|v|a|l|a|t|h|y| 80
Db      3367 GGAGCGGCGCAACGCGCTGAGGAGCTGTGGACACAGCTCAACATGTGGCCCA 3426
Qy      80 |s|h|r|v|a|l|g|l|u|a|a|s|n|a|v|a|l|t|h|y|s|e|r|v|a|l|s|e|r|v|a|l|a|t|h|y| 100
Db      3427 GACCGTGGAGAGGCGGAGAACATCGCGTCACTCGGGGTGGTGGAGGT-GAGCC 3485
Qy      100 |u|a|r|p|r|o|s|e|r| 103
Db      3486 CCGGCCCTCA 3495

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RESULT 15
US-09-949-016-12184
; Sequence 12184, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CLO01307
; CURRENT APPLICATION NUMBER: US/09/949,016

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; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12184
; LENGTH: 8608
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(8608)
; OTHER INFORMATION: n = A,T,C or G
; US-09-949-016-12184

```

```

Alignment Scores:
Pred. No.: 4,696-15 Length: 8608
Score: 201.50 Matches: 97
Percent Similarity: 20.66% Conservative: 3
Best Local Similarity: 20.04% Mismatches: 3
Query Match: 33.03% Indels: 383
DB: Gaps: 2

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US-09-017-715A-2 (1-127) x US-09-949-016-12184 (1-8608)
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Qy      1 |M|e|t|a|s|p|v|a|l|p|h|e|l|y|s|g|l|y|p|h|e|s|e|r|l|e|a|l|y|s|g|l|y|v|a|l|g|l|u 20
Db      2048 ATGATGCTTCAAAAGAGGGCTTCCATCGCAGAGAGGGGTGGTGGTGGTGGTGA 2107
Qy      21 |l|y|t|h|y|s|g|l|y|v|a|l|t|h|r|g|l|u|a|a|a|g|l|y|s|t|h|y|s|g|l|y|v|a|l| 40
Db      2108 AAGACCAAGCAGGGGGGTGCGGAACAGCTGAGAACCAAGAGAGGGGTCAATGATGTG 2167
Qy      40 ----- 40
Db      2168 GGTAAGTGGGCGATGGCAGGGTGGACAGTGTGTGGCCAAAGGTGAGTCCAGTTAC 2227
Qy      40 ----- 40
Db      2228 CTTCGCAAGACTTACTCCCGCAGCCCAAGAGGACATTTGGAGGGGCGAGGCTGG 2287
Qy      40 ----- 40
Db      2288 CTATCAAGGTGGGTCTCCAGACCTTGAGACCAACATGCTGTGCACTATGTG 2347
Qy      40 ----- 40
Db      2348 TGTTTGTCTTTGGCTCTCGGGGCTGTGGGTGTCAGAGACCGAGACAGGCTGGCTA 2407
Qy      40 ----- 40
Db      2408 CCTGCTGTGACAGCACACACATTCCAAGATACAGACCTCCCTGAGGCTGGAGCC 2467
Qy      40 ----- 40
Db      2468 CCTGAAGCCATGAGAGAGCTGTGCTCAGGTGGCCCCACCTCTCCACAGGGAGCGCT 2527
Qy      40 ----- 40
Db      2528 ACAGCGAGTCAAGATCCCTCTCTCCCGAGAGAGAGGGGCAAGGCTGGGGATGAANC 2587
Qy      40 ----- 40
Db      2588 CTAGGCTAGTGTTCCTCCCGCATCTCTCTGACACTCTCCAGAGAGAGAGGGA 2647
Qy      40 ----- 40
Db      2648 GGTCAAGCAATGACTCAGCTCTGGCCCATCTGCTGCTGTTGCTGAGGCCCGGC 2707

```



```

QY      40 ----- 40
Db      2708 CACACCGGGGAGGGGCTGAGACCTGGGTCTAGCCAGTGTCTTACTCTAGAGCTGCTCT 2767
QY      40 ----- 40
Db      2768 CTCTTGTCCCAATCTGTCTGTCCCTTCCATCATCACTTCTTCAGACACAG 2827
QY      40 ----- 40
Db      2828 CAGAGAGAGCCCTGAGAGGGGCGCGCCCAAGACACCATCTTACCCCAACCG 2887
QY      40 ----- 40
Db      2888 ACCCCACAGTTTTCAGAGCTGTCTGTGTGTGTCTGTCTGACCGCCCAACACCTGAG 2947
QY      41 ----- 49
          G1yAla1y8Thr1y8Glu1y8Asn1y8Val1y8
          |||||
Db      2948 GAGAGTCTGGGCTGACAGCTCATTTCTCTCCCAAGAGCCAAAGAGAAATGTTGT 3007
QY      49 1GlnSerVal1ThrSer----- 54
          |||||
Db      3008 ACAGAGCGTGACTTC-AGTGAGAGAGCCAGGGCCAGGGGACATGGGGATAGAGCC 3066
QY      54 ----- 54
Db      3067 CTGGGGCTCCTGCATCTTACTGTCTGGGGCTCAAACTAGAGTCTGCTTACCCCAAC 3126
QY      54 ----- 54
Db      3127 TGGGGTCCCAAGCCCTACAGACCCCTGAGACATGAGGCTAAACTAGGGTGGGCTCTC 3186
QY      54 ----- 54
Db      3187 CTTACCCCAACAGCATCAGAGTGCCCTGAGTCAAGAGGAGCAGGGGTCCAGC 3246
QY      54 ----- 54
Db      3247 AGGGCCAGGGCTTGAGCTCTTGGGAAAGGGCTGCGAGCTGACTCCAGCAGGCTGCT 3306
QY      55 ----- 60
          -----Val1y8Ile1y8Thr1y8
          |||||
Db      3307 TGGGGCTGGGGCTGGGGTGAGAGCCAGCAGTGTCTCCCATAGTGGCCGAGAGACCAA 3366
QY      60 8Glu1y8Ala1y8Asn1y8Val1y8Ser1y8Ala1y8Val1y8Ser1y8Val1y8Asn1y8Thr1y8Val1y8Ala1y8Thr1y8 80
          |||||
Db      3367 GAGAGAGCCCAACCGCTGAGAGGCTGTGTGAGCAGCGCTCAACACTGGGCCACCAA 3426
QY      80 8ThrVal1y8Glu1y8Ile1y8Asn1y8Val1y8Thr1y8Ser1y8Val1y8Ala1y8Arg1y8Glu1y8Asp1y8Le 100
          |||||
Db      3427 GACCGTGGAGAGGCGAGAGACATCGCGTCACTCCGGGTGGTGGCCAGAGT-GAGCC 3485
QY      100 uArgProSer 103
          |||||
Db      3486 CCGGCCCTCA 3495

```

Search completed: May 4, 2005, 09:32:13
 Job time : 305.06 secs

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PI Ji H, Rosen CA;
 XX
 DR MPI, 1998-446811/38.
 DR P-PSDB; AAW63123.
 XX
 XX New isolated human breast cancer specific gene - used to develop products
 PT for the diagnosis, clinical management and treatment of breast cancer and
 PT metastases.
 XX
 PS Claim 4; Fig 1; 73pp; English.

XX This cDNA clone corresponds to the transcript of the newly identified
 CC human breast cancer specific gene 1 (BCSG1), and includes an open reading
 CC frame for a 14.2 kDa protein (see AAW63123). It was isolated from a
 CC breast cancer cDNA library following an EST search for novel genes
 CC differentially expressed in breast cancer versus healthy breast tissue.
 CC The clone is deposited at ATCC 97175 and ATCC 97856. A gradient and stage
 CC -specific BCSG1 expression has been demonstrated from virtually no
 CC detectable expression in normal or benign breast to low level and partial
 CC expression in low grade in situ breast carcinoma and high expression in
 CC infiltrating malignant breast carcinomas. BCSG1 is useful as a breast
 CC cancer progression marker. Recombinant vectors and host cells useful for
 CC recombinant production of BCSG1 polypeptides (including epitope-bearing
 CC polypeptides) are provided. BCSG1 polynucleotides, polypeptides and
 CC antibodies can be used for the detection of breast cancer cells or breast
 CC cancer metastasis, and to develop methods for the clinical management and
 CC treatment of breast cancer
 XX
 SQ Sequence 550 BP; 132 A; 145 C; 192 G; 81 T; 0 U; 0 Other;

Alignment Scores:
 Pred. No.: 6,79e-60 Length: 550
 Score: 610.00 Matches: 127
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 2 Gaps: 0

US-09-017-715A-2 (1-127) x AAW42669 (1-550)

QY 1 Metaspvalphelylsyglpbeserillealalyeglyvalalvalalgin 20
 DB 12 ATGGAATGTTTCAAGAGGGCTTCTCCATCGCCAAAGGGGTGTGGTGGAGAA 71
 QY 21 LysThrlyslngllyvalthrghuilaalagluvalthrlyslngllyvalmettyrval 40
 DB 72 AAGACCAACAGAGGGGTGTGACGAGAGAGCTGAGAAAGCAAGAGGGGTGATGATG 131
 QY 41 GYAlAlAlYsThrlyslngllysluasnvalValGlnSerValThrSerValAlagluysThrlys 60
 DB 132 GGAGCCAAAGACCAAGAGAGATGTTGACAGAGGTGACCTGAGTGCAGAGAACCAAG 191
 QY 61 GIUGlAlAlAsnAlAlValSerlyslalValSerSerValAsnThrValAlathrlys 80
 DB 192 GAGCAGGCCAACGCCCTGAGCAAGGCTGTGTGAGCAGCGTCAACCTGTGGCCACCAAG 251
 QY 81 ThrValGlugluAlagluasnillealValThrSerGlyValValAlArglysluAspLeu 100
 DB 252 ACCGTGAGAGAGCGAGAAACATCGCGTCACTCCGGGGTGTGTGGCAAGGAGCTTG 311
 QY 101 ATGProSerAlAProGlnGlnGlnGlnGlnAlaserlysluysgluGlnuValAlaglu 120
 DB 312 AAGGCATCTGCCCCCAACAGAGAGGTGAGCATCCAAAGAAAGAGAGAGTGGCAGAG 371
 QY 121 GIUAlAGlnSerGlyGlyasp 127
 DB 372 GAGGCCACAGAGTGGGAGAGC 392

RESULT 2
 AAA39470
 ID AAA39470 standard; DNA; 550 BP.
 XX
 AC AAA39470;

XX 24-AUG-2000 (first entry)
 DT
 XX
 DE Human HBGBA67A DNA.
 XX
 KW Human; ADA2; cytosstatic; gene therapy; treatment; cancer;
 XX amyloid-like protein; ss.
 OS Homo sapiens.

XX Key Location/Qualifiers
 FH CDS 12..395
 FT /*tag= a
 FT /product= "HBGBA67"

PN US6054289-A.

PD 25-APR-2000.

PF 30-AUG-1996; 96US-00705771.

PR 30-AUG-1995; 95US-0002993P.

PA (HUMA-) HUMAN GENOME SCI INC.

PI Moore PA;

DR MPI, 2000-338491/29.

DR P-PSDB; AAY87779.

PT New polynucleotide encoding human AD2 is useful for treating cancer and
 PT for isolating cDNAs and genes having similar biological activity.

PS Disclosure; Col 27-28; 54pp; English.

XX This invention describes a novel polynucleotide (I) encoding human ADA2.
 CC The products of the invention have cytosstatic activity and can be used
 CC for gene therapy. (I) is useful for treating cancer; as primers and
 CC probes for isolating full length cDNA and genes having similar biological
 CC activity. This sequence encodes a polypeptide derived from the human
 CC HBGBA67X clone which is an amyloid-like protein found in breast tissue
 XX

SQ Sequence 550 BP; 132 A; 145 C; 192 G; 81 T; 0 U; 0 Other;
 Alignment Scores:
 Pred. No.: 6,79e-60 Length: 550
 Score: 610.00 Matches: 127
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 3 Gaps: 0

US-09-017-715A-2 (1-127) x AAA39470 (1-550)

QY 1 Metaspvalphelylsyglpbeserillealalyeglyvalalvalalgin 20
 DB 12 ATGGAATGTTTCAAGAGGGCTTCTCCATCGCCAAAGGGGTGTGGTGGAGAA 71
 QY 21 LysThrlyslngllyvalthrghuilaalagluvalthrlyslngllyvalmettyrval 40
 DB 72 AAGACCAACAGAGGGGTGTGACGAGAGAGCTGAGAAAGCAAGAGGGGTGATGATG 131
 QY 41 GYAlAlAlYsThrlyslngllysluasnvalValGlnSerValThrSerValAlagluysThrlys 60
 DB 132 GGAGCCAAAGACCAAGAGAGATGTTGACAGAGGTGACCTGAGTGCAGAGAACCAAG 191
 QY 61 GIUGlAlAlAsnAlAlValSerlyslalValSerSerValAsnThrValAlathrlys 80
 DB 192 GAGCAGGCCAACGCCCTGAGCAAGGCTGTGTGAGCAGCGTCAACCTGTGGCCACCAAG 251
 QY 81 ThrValGlugluAlagluasnillealValThrSerGlyValValAlArglysluAspLeu 100
 DB 252 ACCGTGAGAGAGCGAGAAACATCGCGTCACTCCGGGGTGTGTGGCAAGGAGCTTG 311

QY 101 ArgProSerAlaProGlnGlnGlyGluAlaSerIysGluIysGlnGluValAlaGlu 120
DB 312 AGGCGATCTGCCCCCAACAGAGGTGAGGCATCCAAAGAAAGAGAGTGGCAGAG 371
QY 121 GluAlaGlnSerGlyGlyAsp 127
DB 372 GAGGCCCAAGTGTGGGAGAGC 392
RESULT 3
ABLe63343
ID ABL63343 standard; DNA; 550 BP.
XX
AC ABL63343;
XX
DT 15-MAY-2002 (first entry)
XX
DE Breast cancer related gene sequence SEQ ID NO:1680.
XX
KW Human; cancer; colon; breast; ovary; oesophagus; kidney; thyroid;
KW Stomach; lung; prostate; pancreas; carcinoma; antitumour; cancerous;
KW cytostatic; gene therapy; antineoplastic; Wilms' tumour; adenocarcinoma;
KW gene; ds.
XX
OS Homo sapiens.
XX
PN WO200194629-A2.
XX
PD 13-DEC-2001.
XX
PF 30-MAY-2001; 2001WO-US010838.
XX
PR 05-JUN-2000; 2000US-0209473P.
PR 05-JUN-2000; 2000US-0209531P.
PR 18-SEP-2000; 2000US-023133P.
PR 18-SEP-2000; 2000US-023617P.
PR 20-SEP-2000; 2000US-0234009P.
PR 20-SEP-2000; 2000US-0234034P.
PR 20-SEP-2000; 2000US-0234052P.
PR 22-SEP-2000; 2000US-0234509P.
PR 22-SEP-2000; 2000US-0234567P.
PR 25-SEP-2000; 2000US-0234923P.
PR 25-SEP-2000; 2000US-0234924P.
PR 25-SEP-2000; 2000US-0235077P.
PR 25-SEP-2000; 2000US-0235082P.
PR 25-SEP-2000; 2000US-0235134P.
PR 25-SEP-2000; 2000US-0235280P.
PR 26-SEP-2000; 2000US-0235637P.
PR 26-SEP-2000; 2000US-0235638P.
PR 27-SEP-2000; 2000US-0235711P.
PR 27-SEP-2000; 2000US-0235720P.
PR 27-SEP-2000; 2000US-0235840P.
PR 27-SEP-2000; 2000US-0235863P.
PR 28-SEP-2000; 2000US-0236028P.
PR 28-SEP-2000; 2000US-0236032P.
PR 28-SEP-2000; 2000US-0236033P.
PR 28-SEP-2000; 2000US-0236034P.
PR 28-SEP-2000; 2000US-0236109P.
PR 28-SEP-2000; 2000US-0236111P.
PR 29-SEP-2000; 2000US-0236842P.
PR 29-SEP-2000; 2000US-0236891P.
PR 02-OCT-2000; 2000US-0237172P.
PR 02-OCT-2000; 2000US-0237173P.
PR 02-OCT-2000; 2000US-0237278P.
PR 02-OCT-2000; 2000US-0237294P.
PR 02-OCT-2000; 2000US-0237295P.
PR 02-OCT-2000; 2000US-0237316P.
PR 03-OCT-2000; 2000US-0237435P.
PR 03-OCT-2000; 2000US-0237598P.
PR 03-OCT-2000; 2000US-0237604P.
PR 03-OCT-2000; 2000US-0237606P.
PR 03-OCT-2000; 2000US-0237608P.
PR 01-NOV-2000; 2000US-0244867P.

PR 01-NOV-2000; 2000US-0245084P.
XX
XX (AVAL-) AVALON PHARM.
XX
PI Young PE, Augustus M, Carter KC, Ebner R, Endress G, Horrigan S;
PI Soppe DR, Weaver Z;
XX
XX WPI; 2002-188264/24.
DR
XX
PT Screening for anti-neoplastic agent involves exposing cells to a chemical
PT agent to be tested for anti-neoplastic activity, and determining a change
PT in expression of a gene of a signature gene set.
XX
PS Claim 1; SEQ ID NO 1680; 44pp; English.
XX
XX The present invention describes a method (M1) for screening for an anti-
CC neoplastic agent. The method involves exposing cells to a chemical agent
CC to be tested for anti-neoplastic activity, determining a change in
CC expression of at least one gene (I) of a signature gene set, where (I)
CC comprises a sequence (S) selected from 8447 sequences (given in ABL61664
CC to ABL70110), or is at least 95% identical to (S), where a change in
CC expression is indicative of anti-neoplastic activity. (I) has cytostatic
CC activity and can be used in gene therapy. M1 can be used for screening an
CC anti-neoplastic agent, and can be used for producing a product which is
CC the data collected with respect to the anti-neoplastic agent as a result
CC of M1, and the data is sufficient to convey the chemical structure and/or
CC properties of the agent. M1 can be used in the treatment of cancer such
CC as colon, breast, stomach, lung, thyroid, oesophageal, ovarian, kidney,
CC prostate or pancreatic cancer, adenocarcinoma, carcinoma, clear cell
CC cancer, infiltrating ductal cancer, infiltrating lobular cancer, squamous
CC cell carcinoma, neuroendocrine carcinoma, papillary carcinoma and Wilms'
CC tumour
XX
SQ Sequence 550 BP; 132 A; 145 C; 192 G; 81 T; 0 U; 0 Other;
XX
Alignment Scores:
Pred. No.: 6 79e-60 Length: 550
Score: 610.00 Matches: 127
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: Gaps: 0
US-09-017-715A-2 (1-127) x ABL63343 (1-550)
QY 1 MetAspValPheIysIysGlyPheSerIleAlaIysIysGlyValAlaGluValAlaGlu 20
DB 12 ATGAGTGTTCAGAAAGGGCTTCATCGCCAGAAAGGCGGTGGGTGGTGA 71
QY 21 LysThrIysGlnGlyValThrGluAlaAlaGluIysThrIysGluValIleCtyrVal 40
DB 72 AAGACCAAGCAGGGGGTGAAGCAAGCTGCAAGAACCAAGAGGGGTCACTAATG 121
QY 41 GluAlaIysThrIysGluIleValAlaGlnSerValThrSerValAlaGluIysThrIys 60
DB 132 GAGCCCAAGACCAAGAGTGTGTCAGAGCTGAATCTCAATGGCCCAAGACCAAG 191
QY 61 GluGlnAlaAsnAlaValSerIysAlaValIysSerSerValAsnThrValAlaThrIys 80
DB 192 GACGAGGCCAACCCCGTAGCAAGGCTGTAGACGCTCAACACTGTGGCACCAAG 251
QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValAlaIleGlyGluAspLeu 100
DB 252 ACCGTGAGAGAGCGGAGAACATCGGCTCACTCCGGGTGTGTGCCAAGAGGACTTG 311
QY 101 ArgProSerAlaProGlnGlnGlyGluAlaSerIysGluIysGlnGluValAlaGlu 120
DB 312 AGGCGATCTGCCCCCAACAGAGGTGAGGCATCCAAAGAAAGAGAGTGGCAGAG 371
QY 121 GluAlaGlnSerGlyGlyAsp 127
DB 372 GAGGCCCAAGTGTGGGAGAGC 392

```

RESULT 4
ABV73813
ID ABV73813 standard; cDNA; 550 BP.
XX
XX ABV73813;
AC
XX 08-JAN-2003 (first entry)
DT
XX Human gamma-synuclein Glu110 variant gene.
DE
XX Gamma-synuclein; human; single nucleotide polymorphism; SNP;
KM schizophrenia; neuroleptic; gene; ss.
XX
XX Homo sapiens.
OS
FH Key location/Qualifiers
FH CDS 12..395
FT /*tag= a
FT /product= "Gamma-synuclein"
FT /transl_except= (pos:213..215,aa:Glu)
FT replacer(340,T)
FT variation /tag= b
FT /standard_name= "Single nucleotide polymorphism"
FT
XX WO200275317-A2.
XX
XX 26-SEP-2002.
XX
XX 14-MAR-2002; 2002WO-EP002872.
XX
XX 15-MAR-2001; 2001US-0276306P.
XX
XX (NOVS ) NOVARTIS AG.
XX (NOVS ) NOVARTIS-ERFINDUNGEN VERW GES MBH.
XX (UYMA-) UNIV MARYLAND BALTIMORE.
XX
XX Roberts RC, Van Oostrum J, Voshol J, Tamminga CA;
XX
XX WPI: 2002-750574/81.
XX
XX P-PSDB; ABP54932.
XX
XX Screening for compounds for treating or interfering with the onset of
XX Schizophrenia Spectrum Disorders, by detecting interactions of candidate
XX compounds with the gamma-synuclein polypeptide.
XX
XX Disclosure; Fig 1: 32pp; English.
XX
XX The present sequence is that of cDNA encoding the Glu-110 isoform of
XX human gamma-synuclein. The invention relates to an isoform of gamma-
XX synuclein that is caused by an A/T single nucleotide polymorphism (SNP)
XX at position 329 of the gamma-synuclein coding sequence. This SNP causes a
XX glutamic acid to valine change at amino acid position 110 of gamma-
XX synuclein, and is associated with an increased susceptibility of
XX individuals to schizophrenia spectrum disorders (SSDs). This is the first
XX time that a genetic component of SSDs has been identified, and provides a
XX potential target for diagnosis and treatment of schizophrenia. Gamma-
XX synuclein polypeptides, especially those containing the E110V mutation,
XX are used in a claimed method of screening for compounds useful for the
XX treatment of SSDs, and gamma-synuclein expressing cells are used in a
XX claimed method of screening for agonist or antagonist compounds. An
XX oligonucleotide complementary to part of the gamma-synuclein coding
XX sequence is used for the discrimination of an SNP at position 329 of the
XX coding sequence. Gamma-synuclein polypeptides or polynucleotides are also
XX useful for the diagnosis of SSDs, or susceptibility to SSDs, e.g. by PCR
XX amplification of a polynucleotide encoding gamma-synuclein and analysis
XX of the occurrence of the SNP at position 329. A transgenic animal useful
XX for the study of SSDs is also claimed.
XX
XX Sequence 550 BP; 132 A; 145 C; 192 G; 81 T; 0 U; 0 Other;
XX
Alignment Scores: 6.79e-60 Length: 550
Pred. No.: 610.00 Matches: 127
Score:

```

```

Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0
US-09-017-715a-2 (1-127) x ABV73813 (1-550)
QY 1 MetAspValPheIysGlyPheSerIleAlaIysGlyValIglValAlaIglu 20
DB 12 ATGATGTTTTCATAAAGGGGCTTCCTCCATCCGCAAGAGGGGCTGTCGGTGGAA 71
QY 21 LysThrIysGlnGlyValIThrGluAlaIagIulYsThrIysGluIglValMetIYrVal 40
DB 72 AAGACCAAGCAGGGGGGTACGAGACAGCTGAGAGACCAAGAGAGGGGTCTATATGTG 131
QY 41 GIYAlaIysThrIysGluAsnValValIlnSerValIThrSerValAlaIglulYsThrIys 60
DB 132 GGAGCCAAAGACCAAGAGATATTTTACAGACCGTACCTCAGTGGCCGAGAGACCAAG 191
QY 61 GIuGlnAlaAsnAlaValSerIysAlaValIlnSerValIAsnThrValAlaThrIys 80
DB 192 GAGCAGGCGCAACGCCGTGAGCAAGGCTGTGGTGAACAGGTCAACACTGTGGCCACCAAG 251
QY 81 ThrValIgluGluAlaGluAsnIleAlaValIThrSerGlyValValArgIysGluAsnIleu 100
DB 252 ACCGTGAGAGAGCGGAGAACATCCGGTCACTCCGGGGTGTGTGCGAAGAGGACTTGG 311
QY 101 ArgProSerAlaProGlnIgluIgluIlnSerIysGluIysGluValAlaIglu 120
DB 312 AGGCATCTGCCCCCAACAGAGGGGTAGGCATCCAAAGAGAGAAAGAACTGGCAGAG 371
QY 121 GluAlaGlnSerGlyIAsp 127
DB 372 GAGGCCCAAGATCGGGGAGAC 392
RESULT 5
AAD63568
ID AAD63568 standard; cDNA; 550 BP.
XX
XX AAD63568;
AC
XX 12-FEB-2004 (first entry)
DT
XX Human amyloid-like protein cDNA.
DE
XX
XX Human; genetic disease; muscular dystrophy; cystic fibrosis; cytoskeletal;
XX scientific research; gene therapy; gene; amyloid-like protein; ss.
XX
XX Homo sapiens.
XX
XX Key location/Qualifiers
XX FH CDS 12..395
XX FT /*tag= a
XX FT /product= "Human amyloid-like protein"
XX
XX US6639052-B1.
XX
XX 28-OCT-2003.
XX
XX 14-OCT-1999; 99US-00417540.
XX
XX 30-AUG-1995; 95US-0002993P.
XX PR 30-AUG-1996; 96US-00705771.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Moore PA;
XX
XX WPI: 2003-842790/78.
XX
XX P-PSDB; ABW02024.
XX
XX New isolated protein and nucleic acid molecules, useful for diagnostic
XX and therapeutic purposes, e.g. for treating genetic diseases such as
XX

```

PT muscular dystrophy or cystic fibrosis.
 XX
 XX Example 5, Fig 1; Opp, English.
 XX
 CC The invention relates to isolated new isolated protein and nucleic acid
 CC molecules useful for diagnostic and therapeutic purposes. The invention
 CC is for treating genetic diseases such as muscular dystrophy or cystic
 CC fibrosis, and for in vitro purposes related to scientific research,
 CC synthesis of DNA and manufacture of DNA vectors. The invention is useful
 CC in gene therapy. The present sequence is human amyloid-like protein cDNA
 XX
 SQ Sequence 550 BP, 132 A, 145 C, 192 G, 81 T, 0 U, 0 Other;
 Alignment Scores:
 Pred. No.: 6,79e-60 Length: 550
 Score: 610.00 Matches: 127
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 10 Gaps: 0
 US-09-017-715A-2 (1-127) x AAD63568 (1-550)
 QY 1 MetAspValPheLysGlyPheSerIleAlaLysGlyValAlaGluValAlaGlu 20
 DB 12 ATGGATGTTTCAAGAAAGGCTTCTCCATCCCAAGAAAGGCGTGGTGGCGTGA 71
 QY 21 LysThrLysGlnGlyValThrGluAlaAlaGluLysThrLysGlnGlyValMetTyrVal 40
 DB 72 AAGCCAAAGAGGGGGTGCAGCAAGCAAGTGAAGAAAGAGAGGGGTGATGTG 131
 QY 41 GValAlaLysThrLysGluAenValValGlnSerValThrSerValAlaGluLysThrLys 60
 DB 132 GGAAGCCAAAGCAAGAAAGTGTGTACAGAGCTGAGCTCAAGTGGCCGAAGAACCAAG 191
 QY 61 GluGlnAlaAsnAlaValSerLysAlaValAlaSerSerValAenThrValAlaThrLys 80
 DB 192 GAGCAGGCCAAAGCCCGTGAAGCAAGCTGTGTGAGCAAGCCGCAACTGTGGCCACCAAG 251
 QY 81 ThrValGlnGluAlaGluAenIleAlaValThrSerGlyValValAlaGlyGluAspLeu 100
 DB 252 ACCGTGAGAGAGGGGAGAAACATCGCGTCACTCCGGGGTGTGGCCAAAGAGACTTG 311
 QY 101 ArgProSerAlaProGlnGlnGluGluAlaSerLysGluLysGluGluValAlaGlu 120
 DB 312 AGGCCATCTCCGCCCAAGAGAGGTGAGGCATCAAGAGAAAGAGAAAGTGGCAGAG 371
 QY 121 GluAlaGlnSerGlyGlyAsp 127
 DB 372 GAGGCCAGAGTGGGGAGAGAC 392
 RESULT 6
 ADG47636
 ID ADG47636 standard; cDNA; 550 BP.
 XX
 AC ADG47636;
 XX
 DT 11-MAR-2004 (first entry)
 XX
 DE Human amyloid like protein cDNA.
 XX
 XX ss: Gene; muscular dystrophy; cystic fibrosis; hypertension;
 KM angina pectoris; myocardial infarction; ulcer; asthma; allergy;
 KM psychosis; depression; migraine; vomiting; benign prostatic hypertrophy;
 KM osteoporosis; human.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 FT CDS 12..395
 FT /tag= a
 FT /product= "Amyloid like protein"

PN US2003208043-A1.
 XX
 XX 06-NOV-2003.
 XX
 XX 04-JUN-2003; 2003US-00453478.
 XX
 XX 30-AUG-1995; 95US-0002993P.
 XX 30-AUG-1996; 96US-00705771.
 XX 14-OCT-1999; 99US-00417540.
 XX
 XX (HUMA-) HUMAN GENOME SCI INC.
 XX
 PI Moore PA, Gentz RL, Ji H, Ni J, Hu J;
 XX WPI; 2003-864796/80.
 DR P-PSDB; ADG47647.
 XX
 PT New human polypeptides and polynucleotides, useful for diagnosing or
 PT treating genetic diseases such as muscular dystrophy or cystic fibrosis,
 PT hypertension, asthma, depression or osteoporosis.
 XX
 XX Claim 18; SEQ ID NO 1; 56pp; English.
 XX
 CC The invention relates to an isolated human polypeptide. The polypeptides,
 CC polynucleotides, agonists or antagonist are useful for diagnosing or
 CC treating genetic diseases such as muscular dystrophy or cystic fibrosis,
 CC hypertension, angina pectoris, myocardial infarction, ulcers, asthma,
 CC allergies, psychoses, depression, migraine, vomiting, benign prostatic
 CC hypertrophy or osteoporosis. The polypeptides and polynucleotides are
 CC useful for in vitro purposes related to scientific research, synthesis of
 CC DNA and manufacture of DNA vector. The present sequence represents cDNA
 XX encoding human amyloid like protein.
 XX
 SQ Sequence 550 BP, 132 A, 145 C, 192 G, 81 T, 0 U, 0 Other;
 Alignment Scores:
 Pred. No.: 6,79e-60 Length: 550
 Score: 610.00 Matches: 127
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 10 Gaps: 0
 US-09-017-715A-2 (1-127) x ADG47636 (1-550)
 QY 1 MetAspValPheLysGlyPheSerIleAlaLysGlyValAlaGluValAlaGlu 20
 DB 12 ATGGATGTTTCAAGAAAGGCTTCTCCATCCCAAGAAAGGCGTGGTGGCGTGA 71
 QY 21 LysThrLysGlnGlyValThrGluAlaAlaGluLysThrLysGlnGlyValMetTyrVal 40
 DB 72 AAGCCAAAGAGGGGGTGCAGCAAGCAAGTGAAGAAAGAGAGGGGTGATGTG 131
 QY 41 GValAlaLysThrLysGluAenValValGlnSerValThrSerValAlaGluLysThrLys 60
 DB 132 GGAAGCCAAAGCAAGAAAGTGTGTACAGAGCTGAGCTCAAGTGGCCGAAGAACCAAG 191
 QY 61 GluGlnAlaAsnAlaValSerLysAlaValAlaSerSerValAenThrValAlaThrLys 80
 DB 192 GAGCAGGCCAAAGCCCGTGAAGCAAGCTGTGTGAGCAAGCCGCAACTGTGGCCACCAAG 251
 QY 81 ThrValGlnGluAlaGluAenIleAlaValThrSerGlyValValAlaGlyGluAspLeu 100
 DB 252 ACCGTGAGAGAGGGGAGAAACATCGCGTCACTCCGGGGTGTGGCCAAAGAGACTTG 311
 QY 101 ArgProSerAlaProGlnGlnGluGluAlaSerLysGluLysGluGluValAlaGlu 120
 DB 312 AGGCCATCTCCGCCCAAGAGAGGTGAGGCATCAAGAGAAAGAGAAAGTGGCAGAG 371
 QY 121 GluAlaGlnSerGlyGlyAsp 127
 DB 372 GAGGCCAGAGTGGGGAGAGAC 392

ID	ABV73915	standard; cDNA; 550 BP.
XX	ABV73915;	
DT	08-JAN-2003	(first entry)
XX		
DE	Human gamma-synuclein Val110 variant gene.	
XX		
KM	Gamma-synuclein; human; single nucleotide polymorphism; SNP;	
XX	schizophrenia; neuroleptic; mutant; gene; ss.	
OS	Homo sapiens.	
PH		
FT	Key	Location/Qualifiers
FT	CDS	12..395
FT		/tag= a
FT		/product= "Gamma-synuclein"
FT	variation	/transl_except= (pos:213..215,aa:GLU)
FT		replace(340,A)
FT		/tag= b
PN		/standard_name= "Single nucleotide polymorphism"
PD	MO200275317-A2.	
XX	26-SEP-2002.	
XX	14-MAR-2002; 2002MO-EP002872.	
XX	15-MAR-2001; 2001US-0276306P.	
XX		
PA	(NOVS) NOVARTIS AG.	
PA	(NOVS) NOVARTIS-BEFLINDUNGEN VERW GES MBH.	
PA	(UYMA-) UNIV MARYLAND BALTIMORE.	
XX		
PI	Roberts RC, Van Oostrum J, Voshol J, Tamminga CA;	
DR	WPI, 2002-750574/81.	
DR	P-PSDB; ABP54933.	
XX		
PT	Screening for compounds for treating or interfering with the onset of	
PT	Schizophrenia Spectrum Disorders, by detecting interactions of candidate	
PT	compounds with the gamma-synuclein polypeptide.	
XX		
PS	Disclosure; Page; 32pp; English.	
XX		
CC	The present sequence is that of cDNA encoding the Val-110 isoform of	
CC	human gamma-synuclein. The invention relates to an isoform of gamma-	
CC	synuclein that is caused by an A/T single nucleotide polymorphism (SNP)	
CC	at position 329 of the gamma-synuclein coding sequence. This SNP causes a	
CC	glutamic acid to valine change at amino acid position 110 of gamma-	
CC	synuclein, and is associated with an increased susceptibility of	
CC	individuals to schizophrenia spectrum disorders (SSDs). This is the first	
CC	time that a genetic component of SSDs has been identified, and provides a	
CC	potential target for diagnosis and treatment of schizophrenia. Gamma-	
CC	synuclein polypeptides, especially those containing the E110V mutation,	
CC	are used in a claimed method of screening for compounds useful for the	
CC	treatment of SSDs, and gamma-synuclein expressing cells are used in a	
CC	claimed method of screening for agonist or antagonist compounds. An	
CC	oligonucleotide complementary to part of the gamma-synuclein coding	
CC	sequence is used for the discrimination of an SNP at position 329 of the	
CC	coding sequence. Gamma-synuclein polypeptides or polynucleotides are also	
CC	useful for the diagnosis of SSDs, or susceptibility to SSDs, e.g. by PCR	
CC	amplification of a polynucleotide encoding gamma-synuclein and analysis	
CC	of the occurrence of the SNP at position 329. A transgenic animal useful	
CC	for the study of SSDs is also claimed. Note: The present sequence is not	
CC	shown in the specification but is derived from the gamma-synuclein	
CC	sequence given in Fig 1 (see ABV73813)	
XX		
XX	Sequence 550 BP; 131 A; 145 C; 192 G; 82 T; 0 U; 0 Other;	
XX		
XX	Alignment Scores:	

Pred. No.:	4.22e-59	Length:	550
Score:	603.00	Matches:	126
Percent Similarity:	99.21%	Conservative:	0
Best Local Similarity:	99.21%	Mismatches:	1
Query Match:	98.85%	Indels:	0
DB:	6	Gaps:	0

US-09-017-715A-2 (1-127) x ABV73915 (1-550)	
OY 1 MetaspvAlPheLySLyGlyPheSerIleAlLySLyGlyValAlGlyAlaValAlGlu 20	
Db 12 ATGATGTTTTTCAGAAAGAGGGCTTCCTCCATCGCCAAAGAGGGCGTGGTGGTGGGAGAA 71	
OY 21 LysThrLySGInGlyValThrGluAlaAlaGluLysThrLySGInGlyValMetTyVal 40	
Db 72 AAGACCAAGCAGGGGGTACCGAAGCAGCTGGAAGACCAAGAGGGGGTCAATGTTGTG 131	
OY 41 GlyAlaLyThrLySGLuAsnValAlGlnSerValThrSerValAlaGluLysThrLyS 60	
Db 132 GGAGGCCAAGACCAAGAGAAATGTTGTACAGCGTGAACCTCACTGTCGCGAAGACCAAG 191	
OY 61 GluGluAlaAsnAlaValSerLyAlaValSerSerValAsnThrValAlaThrLyS 80	
Db 192 GAGCAGGCCAAGCGCGTGAGCAAGCGTGTGTGACAGCGTCAACACTGTGGCCCAAG 251	
OY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValAlaArgLySGLuAspLeu 100	
Db 252 ACCGTGAGGAGGCGGAGAACATCGCGGTCACTCCGGGGTGTGTGGCAAGAGAGCTTG 311	
OY 101 ArgProSerAlaProGInGInGluGlyGluAlaSerLySGLuLySGluValAlaGlu 120	
Db 312 AGGCCATCTGCCCCCAACAGAGGGGTGTGGCATCCAAAGAAAGAGGAAGTGGCAGAG 371	
OY 121 GluAlaGlnSerGlyGlyAsp 127	
Db 372 GAGGCCAGAGTGGGGAGAGAC 392	

RESULT 8	
AA193778	
ID AA193778 standard; cDNA, 783 BP.	
AC AA193778;	
CC	
DT 06-NOV-2001 (first entry)	
DE Human polynucleotide SEQ ID NO 13838.	
EX	
KW Human; cytokine; cell proliferation; cell differentiation; gene therapy;	
KW tissue growth factor; immunomodulatory; cancer; leukaemia;	
KW nervous system disorders; arthritis; inflammation; ss.	
OS Homo sapiens.	
PN MO200164835-A2.	
PD 07-SEP-2001.	
PF 26-FEB-2001; 2001WO-US004927.	
PR 28-FEB-2000; 2000US-00515126.	
PR 18-MAY-2000; 2000US-00577409.	
PA (HYSE-) HYSEQ INC.	
PI Tang YT, Liu C, Drmanac RT;	
PI WPI; 2001-514838/56.	
DR P-PSDB; AAO13847.	
XX	
PT Isolated nucleic acids and polypeptides, useful for preventing diagnosing	
XX and treating e.g. leukemia, inflammation and immune disorders.	

QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValAlaArgIleGluAspLeu 100
Db 349 ACCGTGAGAGAGCCGAGAAACATCGGTCACCTCCGGGGTGCTGCGCAAGAGACTTG 408
QY 101 ArgProSerAlaProGlnGlnGluGluAlaSerIleGluGluValAlaGlu 120
Db 409 AGGCATCTGCCCCCACCACAGAGGCTGAGCATCCAAAGAAAGAAAGTGGCAGG 468
QY 121 GluAlaGlnSerGlyIleAsp 127
Db 469 GAGGCCACAGATGGGAGAGAC 489
RESULT 10
ADM66887
ID ADM66887 standard; DNA; 488 BP.
XX ADM66887;
AC
XX
XX 03-JUN-2004 (first entry)
DT
XX
XX Human homologue of murine adipocyte specific gamma synuclein DNA Seq 20.
DE
XX
XX human; adipocyte specific; gene; ds; adipose tissue; anti-obesity;
KW high mobility group I-C protein; HMGI-C; obesity; leptin; ob; diabetes;
KW adipogenesis; hypertension; cardiovascular disease; anorectic;
KW antidiabetic; hypotensive; gamma synuclein.
XX
XX Homo sapiens.
OS
XX
XX WO2004011618-A2.
PN
XX
XX 05-FEB-2004.
PD
XX
XX 29-JUL-2003; 2003WO-US023684.
PF
XX
XX 29-JUL-2002; 2002US-0398785P.
PR
XX 12-JUN-2003; 2003US-0478206P.
XX
XX (HMG-) HMG- INC.
PA
XX
XX Chada K, Chouinard R, Ashar H, Sayed AMD;
PI
XX
XX WPI: 2004-143846/14.
DR P-PSDB; ADM67167.
XX
XX Identifying adipocyte specific genes, useful for treating obesity or
PT diabetes, and for identifying drug targets, by differential gene
PT expression analysis between adipose tissue or stromal vascular tissue of
PT mice of different genotypes.
XX
XX Claim 11; SEQ ID NO 20; 91pp; English.
PS
XX
XX This invention relates to a novel method for identifying genes that are
CC over-expressed in adipose tissue and as such it provides targets for anti-
CC -obesity pharmaceutical compositions. Specifically, it refers to a high
CC mobility group I-C protein (HMGI-C) that is associated with obesity and
CC is epistatic to leptin, furthermore, it refers to the ob gene where an
CC autosomal recessive trait is linked to obesity and diabetes. The present
CC invention describes performing differential gene expression analysis
CC between the white adipose tissue (WAT) or stromal vascular tissue (SVT)
CC of any two different mice selected from a group consisting of wild-type,
CC HMGI-C -/-, ob/ob, or HMGI-C -/- ob/ob genotype mice. Accordingly, using
CC this method novel nucleotides and the encoded proteins thereof were
CC identified that are adipocyte specific, and as such can be used for
CC preventing adipogenesis, diagnosing and treating diabetes, obesity,
CC hypertension and cardiovascular disease, as well as screening for
CC compounds that can modulate or prevent adipogenesis and treat diabetes or
CC obesity. These compositions exhibit anorectic, antidiabetic and
CC hypotensive activities. This polynucleotide sequence is a human homologue
CC of a murine adipocyte specific DNA sequence of the invention.
XX
XX Sequence 488 BP; 127 A; 119 C; 176 G; 66 T; 0 U; 0 Other;
SQ

Alignment Scores:
Pred. No.: 2,92e-58 Length: 488
Score: 595.00 Matches: 124
Percent Similarity: 99.21% Conservative: 2
Best Local Similarity: 97.64% Mismatches: 1
Query Match: 97.54% Indels: 0
Gaps: 0
US-09-017-715A-2 (1-127) x ADM66887 (1-488)
QY 1 MetAspValPheIleValSerIleAlaIleValSerGlyValAlaValAlaGlu 20
Db 12 ATGATGCTTTCACAAAGGGCTTCCTCCATCGCCAGAGAGGGGTGTGGATCGGTGGA 71
QY 21 LysThrIleGlnGluValIleThrGluAlaAlaGluIleThrIleGluIleValMetIleVal 40
Db 72 AAGACCAAGCAGGGGGGTGACGAGACAGCTGAGAAAGACCAAGAGGGGTCTATGTGTG 131
QY 41 GluAlaIleThrIleGluAsnValAlaGlnSerValThrSerValAlaGluIleThrIle 60
Db 132 GAGCCCAAGACCAAGAGAGATTTGTACAGACCGTACCTCAGTGGCCGAGAAACCAAG 191
QY 61 GluGlnAlaAsnAlaValSerIleValAlaValSerSerValAsnThrValAlaThrIle 80
Db 192 GAGCAGGCGCAAGCCGTGAGCGAGCTGTGTGAGCAGCAGTCAACACTGTGGCCACCAAG 251
QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValAlaArgIleGluAspLeu 100
Db 252 ACCGTGAGAGAGCCGAGAAACATCGGTCACCTCCGGGGTGCTGCGCAAGAGACTTG 311
QY 101 ArgProSerAlaProGlnGlnGluGluAlaSerIleGluGluValAlaGlu 120
Db 312 AGGCATCTGCCCCCACCACAGAGGCTGAGCATCCAAAGAAAGAAAGTGGCAGG 371
QY 121 GluAlaGlnSerGlyIleAsp 127
Db 372 GAGGCCACAGATGGGAGAGAC 392
RESULT 11
AAK29997
ID AAK29997 standard; DNA; 720 BP.
XX
XX AAK29997;
AC
XX
XX 06-JUL-1999 (first entry)
DT
XX
XX Human peryon gene.
DE
XX
XX Human; synuclein; peryon; diagnosis; neurodegenerative disorder; cancer;
KW breast; skin; intermediate filament damage; ss.
XX
XX
XX Homo sapiens.
OS
XX
XX EP908727-A1.
PN
XX
XX 14-APR-1999.
PD
XX
XX 21-SEP-1998; 98EP-00307628.
PF
XX
XX 19-SEP-1997; 97GB-00019879.
PR
XX
XX (NEUR-) NEUROPA LTD.
PA (UYSA-) UNIV ST ANDREWS.
XX
XX WPI: 1999-217169/19.
DR P-PSDB; AAY07271.
XX
XX New synuclein protein (peryon) and gene, useful in assays for screening,
PT diagnosing or monitoring cancer, neurodegenerative disorders or skin
PT disorders.
XX
XX Claim 29; Page 16-17; 39pp; English.
PS

DB 169 GGAGCCAGACAGCAGAGATGTTGTACAGAGCGTGACCTGACGGCCGAGAAAGCAAG 228
QY 61 GUGGUAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAG 80
DB 229 GAGCAGGCGCAGAGCGGCGGAGCGAGCGGCGGAGCGGAGCGGAGCGGAGCGGAG 288
QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValAlaArgGlyGluAspLeu 100
DB 289 ACCGTGGAGAGAGCGAGAAACATCGCGGTCACTCCGGGGTGGTGGCAAGAGAGACTTG 348
QY 101 ArgProSerAlaProGlnGlnGluGluAlaSerGlyGluGlyGluValAlaGlu 120
DB 349 AGCCCATCTGCCCGCCCAACAGAGGGGTGTGCATCTCAAGAGAAAGAGAGAGTGCAGAG 408
QY 121 GluAlaGlnSerGlyGlyAsp 127
DB 409 GAGGCCAGAGTGGGGAGAGC 429
RESULT 13
ADE43864
ID ADE43864 standard; cDNA; 720 BP.
XX ADE43864;
XX 29-JAN-2004 (first entry)
DE Human SNCG cDNA, SEQ ID 469.
XX
XX Neurodegenerative disease; uPA; SNCG; IDE; KNSL1; LIPA; TNFRSF6;
KW Alzheimer's disease; neuroprotective; nootropic; gene therapy;
XX Chromosome 10; gene; ss.
XX
XX Homo sapiens.
XX WO2003054143-A2.
XX
XX 03-JUL-2003.
XX 25-OCT-2002; 2002WO-US034679.
XX
XX 25-OCT-2001; 2001US-0339525P.
PR 08-NOV-2001; 2001US-0336929P.
PR 08-NOV-2001; 2001US-0338010P.
PR 09-NOV-2001; 2001US-0338363P.
PR 04-DEC-2001; 2001US-0337052P.
PR 28-MAR-2002; 2002US-0368919P.
XX
XX (NEUR-) NEUROGENETICS INC.
PA (GEHO) GEN HOSPITAL CORP.
XX
PI Becker KD, Veliceljebi G, Elliott KJ, Wang X, Tanzi RE, Bertram L;
PI Saunders AJ, Mullin KM, Sampson AJ, Blacker DL;
XX
XX WPI; 2003-559131/52.
XX
XX Determining a predisposition for or the occurrence of neurodegenerative
PT disease, e.g. Alzheimer's disease by detecting in a target nucleic acid
PT the presence or absence of an allelic variant of one or more polymorphic
PT regions.
XX
XX Claim 84; Page 740; 848p; English.
XX
XX The present invention relates to a method (M1) for determining a
XX predisposition for or the occurrence of neurodegenerative disease in a
XX subject. The method comprises detecting in a target nucleic acid obtained
XX from the subject the presence or absence of an allelic variant of one or
XX more polymorphic regions of one or more genes selected from uPA
XX (urokinase plasminogen activator), SNCG (gamma-synuclein), IDE (insulin-
XX degrading enzyme), KNSL1 (Kinesin-like protein 1), LIPA (lysosomal acid
XX lipase), and TNFRSF6 (Tumour Necrosis Factor Receptor-SF6), where the
XX presence of at least one of the allelic variant of one or more
XX polymorphic regions is indicative of a predisposition for or the
XX occurrence of neurodegenerative disease. The genes are all located on

CC chromosome 10. M1 is useful for determining a predisposition for or the
CC occurrence of, and for treating neurodegenerative disease, particularly
CC Alzheimer's disease.
XX
XX SQ Sequence 720 BP; 172 A; 205 C; 212 G; 120 T; 0 U; 11 Other;
XX
XX Alignment Scores:
XX Pred. No.: 1,056-57 Length: 720
XX Score: 592.00 Matches: 124
XX Percent Similarity: 97.64% Conservative: 0
XX Best Local Similarity: 97.64% Mismatches: 3
XX Query Match: 97.05% Indels: 0
XX DB: 10 Gaps: 0
XX
XX US-09-017-715a-2 (1-127) x ADE43864 (1-720)
QY 1 MetAspValPheLysGlyPheSerIleAlaLysGlyValAlaGlyValAlaGlu 20
DB 49 ATGATGTTTTCAGAAAGGGCTTCTCCATCCGCAAGNAGGCGGTGGTGGCGGAGAA 108
QY 21 LysThrLysGlnGlyValThrGluAlaGluLysThrLysGlnGlyValMetTyVal 40
DB 109 AAGACCAAGCAGGCGGTGACGAGACAGCTGAGAAAGACCAGAGAGGGGTCAATGATG 168
QY 41 GlyAlaLysThrLysGluAsnValAlaGlnSerValThrSerValAlaGluLysThrLys 60
DB 169 GGAGCCAGACCAAGAGAGATGTTGTACAGACGTCGACCTGCTGTCGCGAGAAAGCAAG 228
QY 61 GUGGUAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAG 80
DB 229 GAGCAGGCGCAGAGCGGCGGAGCGGAGCGGAGCGGAGCGGAGCGGAGCGGAG 288
QY 81 ThrValGluGluAlaGluAsnIleAlaValThrSerGlyValAlaArgGlyGluAspLeu 100
DB 289 ACCGTGGAGAGAGCGAGAAACATCGCGGTCACTCCGGGGTGGTGGCAAGAGAGACTTG 348
QY 101 ArgProSerAlaProGlnGlnGluGluAlaSerGlyGluGlyGluValAlaGlu 120
DB 349 AGCCCATCTGCCCGCCCAACAGAGGGGTGTGCATCTCAAGAGAAAGAGAGAGTGCAGAG 408
QY 121 GluAlaGlnSerGlyGlyAsp 127
DB 409 GAGGCCAGAGTGGGGAGAGC 429
RESULT 14
ADH54342
ID ADH54342 standard; cDNA; 720 BP.
XX
XX ADH54342;
XX 25-MAR-2004 (first entry)
DE Human SNCG gene cDNA sequence SeqID469.
XX
XX human; neurodegenerative disease; urokinase plasminogen activator; uPA;
KW gamma-synuclein; SNCG; insulin degrading enzyme; IDE;
KW kinesin-like protein 1; KNSL1; lysosomal acid lipase; LIPA;
KW tumour necrosis factor receptor SF6; TNFRSF6; Alzheimer's disease; ss.
XX
XX Homo sapiens.
XX
XX US2003224380-A1.
XX
XX 04-DEC-2003.
XX 25-OCT-2002; 2002US-00282174.
XX
XX 25-OCT-2001; 2001US-0339525P.
PR 25-OCT-2001; 2001US-0348065P.
PR 02-NOV-2001; 2001US-0336983P.
PR 08-NOV-2001; 2001US-0336929P.
PR 08-NOV-2001; 2001US-0338010P.
PR 09-NOV-2001; 2001US-0338363P.

PR 04-DEC-2001; 2001US-0337052P.
PR 28-MAR-2002; 2002US-0368919P.
XX (GEHO) GEN HOSPITAL CORP.
XX Becker KD, Velicelbi G, Elliott KJ, Wang X, Tanzi RE;
PI Bertram L, Saunders AJ, Mullin KM, Sampson AJ;
XX WPI; 2004-060538/06.
XX
XX Determining a predisposition for or the occurrence of neurodegenerative
PT disease, particularly Alzheimer's disease, comprises determining the
PT presence of a polymorphism in the UPA, SNCG, IDE, KNSL1, LIPA or TNFRSF6
PT gene.
XX
XX Claim 84; SEQ ID NO 469; 205pp; English.
XX
XX This invention relates to a novel method of determining a predisposition
CC for or the occurrence of neurodegenerative disease comprising detecting
CC in a target nucleic acid obtained from the subject the presence of an
CC allelic variant of polymorphic regions of human genes selected from
CC urokinase plasminogen activator (uPA), gamma-synuclein (SNCG), insulin
CC degrading enzyme (IDE), kinesin-like protein 1 (KNSL1), lysosomal acid
CC lipase (LIPA) and tumour necrosis factor receptor SF6 (TNFRSF6). The
CC method is useful in determining the presence or predisposition to a
CC neurodegenerative disease, particularly Alzheimer's disease. The present
CC sequence is the cDNA sequence of the human SNCG gene which is related to
CC the invention.
XX
XX Sequence 720 BP; 172 A; 205 C; 212 G; 120 T; 0 U; 11 Other;
SQ
XX
XX Alignment Scores:
Pred. No.: 1.05e-57 Length: 720
Score: 592.00 Matches: 124
Percent Similarity: 97.64% Conservative: 0
Best Local Similarity: 97.64% Mismatches: 3
Query Match: 97.05% Indels: 0
Gaps: 0
DB: 12
US-09-017-715A-2 (1-127) x ADH54342 (1-720)
QY 1 MetAspValPheIySGlyPheSerIeAlaIySGlyValIaGlyAlaIaGlu 20
Db 49 ATGATGTTTCAAGAGGCGTTCTCATGCCAAGAGGCGGTGGTGGCGTGA 108
QY 21 LysThrIySGlnGlyValIThrGluAlaIaGluIyThrIySGlnGlyValI 40
Db 109 AAGACCAAGGAGGGGTGACGGAAGCAGCTGAGAAAGCAAGGAGGGGTATGTG 168
QY 41 GAlaIaIyThrIySGlnGluIyValIaGlnSerIyThrIySGlnGlyValI 60
Db 169 GAGGCCAAGGCAAGGAGGAAATGTTGTCAGAGCGTGAAGGCGCGGCAAGCAAG 228
QY 61 GAlaIa 80
Db 229 GAGCAGGCGCAACGCGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 288
QY 81 ThrValIaGlnGluIaGlnIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 100
Db 289 ACCGTGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 348
QY 101 ArgProSerIa 120
Db 349 AGGCCATCTCTCCCGCCCAAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 408
QY 121 GluAlaGlnSerIyGlyIaSP 127
Db 409 GAGGCCAGAGTGGGAGGAGC 429
RESULT 15
ACH15493
XX ACH15493 standard; cDNA; 479 BP.
XX

AC ACH15493;
XX 13-OCT-2003 (first entry)
XX Human adult brain cDNA #2705.
XX
XX Human; seq; sequencing by hybridisation; SBH; expressed sequence tag; EST;
XX genome mapping; biodiversity; genetic disorder.
XX
XX Homo sapiens.
XX
XX US2003073623-A1.
XX
XX 17-APR-2003.
XX
XX 30-JUL-2001; 2001US-00918995.
XX
XX 30-JUL-2001; 2001US-00918995.
XX
XX 30-JUL-2001; 2001US-00918995.
XX
XX (DRMA/) DRMANAC R T.
XX (LABA/) LABAT I.
XX (STAC/) STACHE-CRAIN B.
XX (DICK/) DICKSON M C.
XX (JONE/) JONES L W.
XX
XX Drmanac RT, Labat I, Stache-Crain B, Dickson MC, Jones LW;
XX WPI; 2003-615964/58.
XX
XX New polynucleotide sequences obtained from various cDNA libraries, useful
PT as hybridization probes, as oligomers for PCR, for chromosome and gene
PT mapping, in the recombinant production of protein, or in generating
PT antisense DNA or RNA.
XX
XX Claim 1; SEQ ID NO 2705; 44pp; English.
XX
XX The invention relates to an isolated polynucleotide comprising any one of
CC 38043 cDNA sequences, appearing as ACH12789-ACH50831, whose sequence was
CC determined by the technique of SBH (sequencing by hybridisation). Also
CC included is a purified polypeptide comprising a sequence corresponding to
CC a reading frame of the novel polynucleotide. The nucleic acid sequences
CC are useful in diagnostics as expressed sequence tags (EST) for
CC identifying expressed genes or for physical mapping of the human genome,
CC in forensics, in assessing biodiversity, or in identifying mutations,
CC responsible for genetic disorders and other traits. The nucleotide
CC sequences are also useful as hybridisation probes, as oligomers for PCR,
CC for chromosome and gene mapping, in the recombinant production of
CC protein, or in generating antisense DNA or RNA. The purified polypeptide
CC is useful for generating antibodies specific for it. The present sequence
CC is one of the 38043 isolated cDNA/EST sequences. Note: The sequence data
CC for this patent did not form part of the printed specification, but was
CC obtained in electronic format directly from USPTO at
CC seqdata.uspto.gov/sequence.html?docid=20030073623
XX
SQ Sequence 479 BP; 120 A; 118 C; 178 G; 56 T; 0 U; 7 Other;
SQ
XX
XX Alignment Scores:
Pred. No.: 8.23e-52 Length: 479
Score: 538.00 Matches: 112
Percent Similarity: 99.14% Conservative: 3
Best Local Similarity: 96.55% Mismatches: 1
Query Match: 88.20% Indels: 0
Gaps: 0
DB: 9
US-09-017-715A-2 (1-127) x ACH15493 (1-479)
QY 12 LysIySGlnGlyValIaGlnIyThrIySGlnGlyValIaGlnIyThrIySGln 31
Db 47 CAGGAGGCGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 106
QY 32 LysThrIySGlnGlyValIaGlnIyThrIySGlnGlyValIaGlnIyThrIySGln 51
Db 107 AAGACCAAGGAGGGGTGATGTGTGTGGAGCCCAAGGAGGAGGAGGAGGAGGAG 166

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OY 52 ValThrSerValAlaGluIysThrIysGluGlnAlaAsnAlaValSerIysAlaValAla 71
Db 167 GTGACCTCAGTGGCCGAGAGACCAAGAGCCAGCCGCTGAGCGAGGCTGTGGTG 226
OY 72 SerSerValAsnThrValAlaThrIysThrValGluGluAlaGluAsnIleAlaValThr 91
Db 227 AGCAGCGTCACACACTGTGGCCACCAAGACCCTGTGAGAGAGCGGAGAACATCGCGGTCAAC 286
OY 92 SerGlyValIvalArgIysGluAspLeuArgProSerAlaProGlnGlnGluGluAla 111
Db 287 TCCGGGGGTGTGGCAGAGGAGCACTTGAGGCCATCTGCCCCCAACAGAGGGGTGAGGCA 346
OY 112 SerIysGluIysGluGluValAlaGluGluAlaGlnSerGlyValAsp 127
Db 347 TCCAAAGAGAAAGAGAGAGTGGCAGAGAGGCCCAAGAGTGGGGGAGAGC 394
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